

U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE

BARNETT'S DOCKET NUMBER

68 USOP

TRANSMITTAL LETTER TO THE UNITED STATES
DESIGNATED/ELECTED OFFICE (DO/EO/US)
CONCERNING A FILING UNDER 35 U.S.C. 371

U.S. APPLICATION NO. (If known, see 37 CFR 1.5)

10/088771

INTERNATIONAL APPLICATION NO.
PCT/JP00/06375

INTERNATIONAL FILING DATE
September 19, 2000

PRIORITY DATE CLAIMED
September 20, 1999

TITLE OF INVENTION

Melanin Concentrating Hormone Antagonist

APPLICANT(S) FOR DO/EO/US

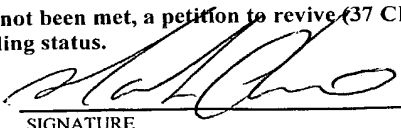
Kaneyoshi KATO, Jun TERAUCHI, Masaaki MORI, Nobuhiro SUZUKI, Yukio SHIMOMURA, Shiro TAKEKAWA, Yuji ISHIHARA

Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:

1. ☒ This is a **FIRST** submission of items concerning a filing under 35 U.S.C. 371.
2. ☐ This is a **SECOND** or **SUBSEQUENT** submission of items concerning a filing under 35 U.S.C. 371.
3. ☐ This is an express request to begin national examination procedures (35 U.S.C. 371(f)). The submission must include items (5), (6), (9) and (21) indicated below.
4. ☒ The US has been elected by the expiration of 19 months from the priority date (Article 31).
5. ☒ A copy of the International Application as filed (35 U.S.C. 371(c)(2))
 - a. ☐ is attached hereto (required only if not communicated by the International Bureau).
 - b. ☒ has been communicated by the International Bureau.
 - c. ☐ is not required, as the application was filed in the United States Receiving Office (RO/US).
6. ☐ An English language translation of the International Application as filed (35 U.S.C. 371(c)(2)).
 - a. ☒ is attached hereto.
 - b. ☐ has been previously submitted under 35 U.S.C. 154(d)(4).
7. ☐ Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3))
 - a. ☒ are attached hereto (required only if not communicated by the International Bureau).
 - b. ☐ have been communicated by the International Bureau.
 - c. ☐ have not been made; however, the time limit for making such amendments has NOT expired.
 - d. ☐ have not been made and will not be made.
8. ☐ An English language translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371 (c)(3)).
9. ☒ An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)).
10. ☐ An English language translation of the annexes of the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371(c)(5)).

Items 11 to 20 below concern document(s) or information included:

11. ☒ An Information Disclosure Statement under 37 CFR 1.97 and 1.98.
12. ☒ An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.
13. ☒ A FIRST preliminary amendment.
14. ☐ A SECOND or SUBSEQUENT preliminary amendment.
15. ☐ A substitute specification.
16. ☐ A change of power of attorney and/or address letter.
17. ☒ A computer-readable form of the sequence listing in accordance with PCT Rule 13ter.2 and 35 U.S.C. 1.821 - 1.825.
18. ☐ A second copy of the published international application under 35 U.S.C. 154(d)(4).
19. ☐ A second copy of the English language translation of the international application under 35 U.S.C. 154(d)(4).
20. ☒ Other items or information:
Itemized Return Postcard; Certificate of Express Mailing; Express Mail Label No. EL 916492917 US
PCT Forms 101, 301, 304, 308, 332; Written Opinion; Date of Deposit March 19, 2002
Recordation Cover Sheet; Cited References (15)

U.S. APPLICATION NO. (if known, see 37 CFR 1.53) 10/088771		INTERNATIONAL APPLICATION NO. PCT/JP00/06375		ATTORNEY'S DOCKET NUMBER 2648 USOP			
21. <input checked="" type="checkbox"/> The following fees are submitted: BASIC NATIONAL FEE (37 CFR 1.492 (a) (1) - (5)): Neither international preliminary examination fee (37 CFR 1.482) nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO and International Search Report not prepared by the EPO or JPO. \$1040.00 International preliminary examination fee (37 CFR 1.482) not paid to USPTO but International Search Report prepared by the EPO or JPO \$890.00 International preliminary examination fee (37 CFR 1.482) not paid to USPTO but international search fee (37 CFR 1.445(a)(2)) paid to USPTO \$740.00 International preliminary examination fee (37 CFR 1.482) paid to USPTO but all claims did not satisfy provisions of PCT Article 33(1)-(4) \$710.00 International preliminary examination fee (37 CFR 1.482) paid to USPTO and all claims satisfied provisions of PCT Article 33(1)-(4) \$100.00 ENTER APPROPRIATE BASIC FEE AMOUNT =				CALCULATIONS PTO USE ONLY <table border="1" style="width:100%; border-collapse: collapse;"> <tr> <td style="width:50%; text-align: right;">\$ 890.00</td> <td style="width:50%;"></td> </tr> </table>		\$ 890.00	
				\$ 890.00			
Surcharge of \$130.00 for furnishing the oath or declaration later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492(e)).							
CLAIMS	NUMBER FILED	NUMBER EXTRA	RATE	\$			
Total claims	33 - 20 =	13	x \$18.00	\$	234.00		
Independent claims	10 - 3 =	7	x \$84.00	\$	588.00		
MULTIPLE DEPENDENT CLAIM(S) (if applicable)			+ \$280.00	\$			
TOTAL OF ABOVE CALCULATIONS =				\$			
<input type="checkbox"/> Applicant claims small entity status. See 37 CFR 1.27. The fees indicated above are reduced by 1/2.				\$			
SUBTOTAL =				\$	1,712.00		
Processing fee of \$130.00 for furnishing the English translation later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492(f)).				\$			
TOTAL NATIONAL FEE =				\$	1,712.00		
Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31). \$40.00 per property +				\$			
TOTAL FEES ENCLOSED =				\$	1,712.00		
				Amount to be refunded:	\$		
				charged:	\$		
a. <input type="checkbox"/> A check in the amount of \$ _____ to cover the above fees is enclosed. b. <input checked="" type="checkbox"/> Please charge my Deposit Account No. <u>500799</u> in the amount of \$ <u>1,712.00</u> to cover the above fees. A duplicate copy of this sheet is enclosed. c. <input checked="" type="checkbox"/> The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any overpayment to Deposit Account No. <u>500799</u> . A duplicate copy of this sheet is enclosed. d. <input type="checkbox"/> Fees are to be charged to a credit card. WARNING: Information on this form may become public. Credit card information should not be included on this form. Provide credit card information and authorization on PTO-2038.							
NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137 (a) or (b)) must be filed and granted to restore the application to pending status.							
SEND ALL CORRESPONDENCE TO: Mark Chao, PhD, JD Takeda Pharmaceuticals North America, Inc. Suite 500, 475 Half Day Road Lincolnshire, IL 60069 USA (847)383-3372 fax (847)383-3481							
				 SIGNATURE Mark Chao, PhD, JD NAME 37,293 REGISTRATION NUMBER For Customer No. 23,115			

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Application No.: National Phase of PCT/JP00/06375
Filed: tba
1st Inventor: KATO, Kaneyoshi
For: Melanin Concentrating Hormone
Antagonist
Atty. Dkt. No. 2648 USOP

Art Unit: tba
Examiner: tba
Allowed:
Batch:
Paper No.:

#4/a

Preliminary Amendment

BOX NEW APPLICATION
Assistant Commissioner for Patents
Washington, D.C. 20231
Sir:

AMENDMENT

In the Specification

Please insert on Page 1 as the first sentence of the application the following:

-- This application is the National Phase filing of International
Patent Application No. PCT/JP00/06375, filed September 19,
2000. --

In the Claims

Please Cancel Claims 3, 13, 21, 24, 27, 29, 30, 31, 32, 33, 37 and 38, without prejudice to
future continuing applications.

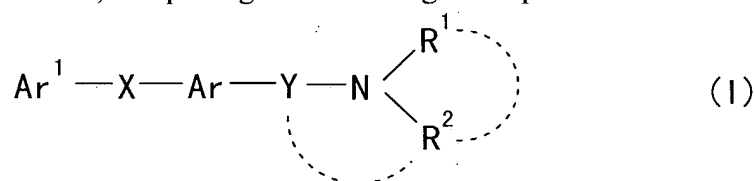
Please add new claims 39-45 (inclusive) as shown below.

Please amend claim 17 to read as shown below.

A mark-up of the claims showing the requested change also follows.

17. (AMENDED) A pharmaceutical composition, which comprises a melanin-concentrating hormone antagonist in combination with at least one species selected from the group consisting of an agent for treating diabetes, an agent for treating hypertension and an agent for treating arteriosclerosis and a pharmaceutically acceptable carrier, diluent or excipient.

39. (NEW) A method for antagonizing melanin-concentrating hormone in a mammal in need thereof, comprising administering a compound of the formula :



wherein Ar^1 is a cyclic group which may have substituents;

X is a spacer having a main chain of 1 to 6 atoms;

Y is a bond or a spacer having a main chain of 1 to 6 atoms;

Ar is a monocyclic aromatic ring which may be condensed with a 4 to 8 membered non-aromatic ring, and may have further substituents;

R^1 and R^2 are independently hydrogen atom or a hydrocarbon group which may have substituents; R^1 and R^2 , together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; R^2 may form a spiro ring together with Ar; or R^2 , together with the adjacent nitrogen atom and Y, may form a nitrogen-containing hetero ring which may have substituents; or a salt thereof to a mammal.

40. (NEW) A pharmaceutical composition which comprises a compound as defined in claim 18 and a pharmaceutically acceptable carrier, diluent or excipient.

41. (NEW) A pharmaceutical composition which comprises a compound as defined in claim 19, and a pharmaceutically acceptable carrier, diluent or excipient.

42. (NEW) A pharmaceutical composition which comprises a compound as defined in claim 22 and a pharmaceutically acceptable carrier, diluent or excipient.

43. (NEW) A pharmaceutical composition which comprises a compound as defined in claim 25 and a pharmaceutically acceptable carrier, diluent or excipient.

44. (NEW) A pharmaceutical composition which comprises a compound as defined in claim 26 and a pharmaceutically acceptable carrier, diluent or excipient.

45. (NEW) A pharmaceutical composition which comprises a compound as defined in claim 28 and a pharmaceutically acceptable carrier, diluent or excipient.

Mark-up of the claims showing changes.

17. (AMENDED) A pharmaceutical composition, which comprises a melanin-concentrating hormone antagonist in combination with at least one species selected from the group consisting of an agent for treating diabetes, an agent for treating hypertension and an agent for treating arteriosclerosis and a pharmaceutically acceptable carrier, diluent or excipient.

REMARKS

The specification is amended above to insert a reference to related cases.

The claims have been amended to better conform the claims to accepted U.S. practice.

Support for amendment to claim 17 can be found in the specification as filed.

Claim 39 encompasses the subject matter of cancelled claims 37 and 38, and is supported by the specification as filed.

Claims 40-45 encompass the subject matter of cancelled claim 32, rewritten in dependent form.

No amendment of inventorship is necessitated by these amendments.

A clean copy of the claims as amended are attached for the Examiner's use.

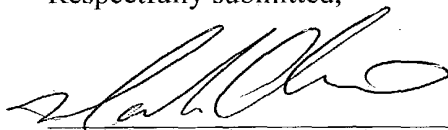
Early allowance of the claims is requested. Should the Examiner believe that a conference with applicants' attorney would advance prosecution of this application, the Examiner is respectfully invited to call applicants' attorney.

Respectfully submitted,

Dated: March 18, 2002

(847)383-3372

(847)383-3391



Mark Chao, Ph.D., Reg. No. 37,293

Elaine M. Ramesh, Ph.D., Reg. No. 43032

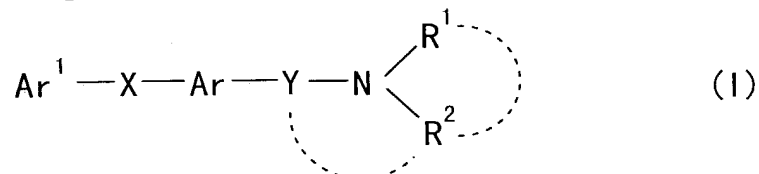
Attorney for Applicants

Customer No. 23115

Takeda Pharmaceuticals North America, Inc.
Intellectual Property Department
Suite 500, 475 Half Day Road
Lincolnshire, IL 60069 USA

CLAIMS

1. A melanin-concentrating hormone antagonist which comprises a compound of the formula :



5 wherein Ar¹ is a cyclic group which may have substituents;
 X is a spacer having a main chain of 1 to 6 atoms;
 Y is a bond or a spacer having a main chain of 1 to 6
 10 atoms;
 Ar is a monocyclic aromatic ring which may be condensed with a 4 to 8 membered non-aromatic ring, and may have further substituents;
 R¹ and R² are independently hydrogen atom or a
 15 hydrocarbon group which may have substituents; R¹ and R², together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; R² may form a spiro ring together with Ar; or R², together with the adjacent nitrogen atom and
 20 Y, may form a nitrogen-containing hetero ring which may have substituents; or a salt thereof.

2. An antagonist according to claim 1, wherein Y is a
 25 spacer having a main chain of 1 to 6 atoms; R¹ and R² are independently hydrogen atom or a hydrocarbon group which may have substituents; R¹ and R², together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; or R² may form a spiro ring together with Ar.

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3. (Cancelled)

4. An antagonist according to claim 1, wherein the cyclic group for Ar^1 is C_{6-14} monocyclic or condensed polycyclic aromatic hydrocarbon group.
5. An antagonist according to claim 1, wherein the cyclic group for Ar^1 is a group formed by removing an optional one hydrogen atom from an aromatic ring assemble in which 2 or 3 C_{6-14} monocyclic or condensed polycyclic aromatic hydrocarbon groups are directly bonded by single bonds.
6. An antagonist according to claim 1, wherein the cyclic group for Ar^1 is a group formed by removing an optional one hydrogen atom from an aromatic ring assemble in which C_{6-14} monocyclic or condensed polycyclic aromatic hydrocarbon and 5 to 10 membered aromatic hetero ring are directly bonded by a single bond.
7. An antagonist according to claim 1, wherein Ar^1 is phenyl, biphenyl, phenyl-pyridyl, phenyl-furyl, phenyl-isoxazolyl, diphenyl-oxazolyl, pyridyl-phenyl, phenyl-pyrimidinyl, benzofuranyl-phenyl, furyl-phenyl, terphenyl, thienyl-phenyl, indolyl, naphthyl-oxadiazolyl, benzofuranyl-oxadiazolyl, benzothienyl, benzofuranyl, fluorenyl, pyridyl-pyrrolyl or thioxanthanyl; each of which may have 1 to 3 substituents selected from the group consisting of halogen atom; nitro; C_{1-3} alkylenedioxy; optionally halogenated C_{1-6} alkyl; hydroxy- C_{1-6} alkyl; optionally halogenated C_{3-6} cycloalkyl; optionally halogenated C_{1-6} alkoxy; optionally halogenated C_{1-6} alkythio; hydroxy; C_{7-19} aralkyloxy which may have substituents; C_{6-14} aryloxy which may have substituents; amino; mono- C_{1-6}

alkylamino; di-C₁₋₆ alkylamino; 5 to 7 membered saturated cyclic amino which may have substituents and may be condensed with a benzene ring; 5 to 7 membered non-aromatic heterocyclic groups which may have substituents; formyl; carboxy; C₆₋₁₄ aryl-carbonyl which may have substituents; C₆₋₁₄ aryl-carbamoyl which may have substituents; aromatic hetero ring-carbamoyl which may have substituents; C₁₋₆ alkoxy-carbonyl; optionally halogenated C₁₋₆ alkyl-carboxamide; C₆₋₁₄ aryl-carboxamide which may have substituents; C₇₋₁₉ aralkyl-carboxamide which may have substituents; aromatic hetero ring-carboxamide which may have substituents; N-(C₆₋₁₄ aryl-carbonyl which may have substituents)-N-C₁₋₆ alkylamino; C₆₋₁₄ arylamino-carbonylamino which may have substituents; C₆₋₁₄ arylsulfonylamino which may have substituents; C₆₋₁₄ aryl-carbonyloxy which may have substituents; oxo; carboxy-C₁₋₆ alkyl; C₁₋₆ alkoxy-carbonyl-C₁₋₆ alkyl; C₇₋₁₉ aralkyl which may have substituents; aromatic hetero ring-C₁₋₆ alkoxy; and cyano.

8. An antagonist according to claim 1, wherein Ar¹ is piperidinyl, piperazinyl, pyrrolidinyl, dihydropyridyl or tetrahydropyridyl; each of which may have 1 or 2 substituents selected from the group consisting of oxo, C₆₋₁₄ aryl which may have substituents, hydroxy, C₇₋₁₉ aralkyloxy-carbonyl, and C₇₋₁₉ aralkyl.

9. An antagonist according to claim 1, wherein the "spacer having a main chain of 1 to 6 atoms" for X and Y is a bivalent group consisting of 1 to 3 species selected from -O-, -S-, -CO-, -SO-, -SO₂-, -NR⁸- (R⁸ is hydrogen atom, optionally halogenated C₁₋₆ alkyl, optionally halogenated C₁₋₆ alkyl-carbonyl, optionally halogenated C₁₋₆ alkylsulfonyl), and a bivalent C₁₋₆ non-

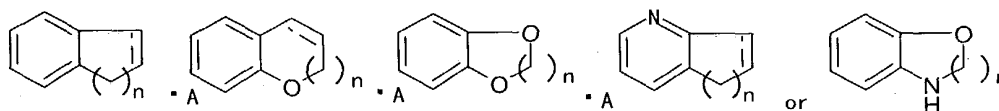
cyclic hydrocarbon group which may have substituents.

10. An antagonist according to claim 1, wherein X is -
 CONR^{8c}-, -NR^{8c}CO-, -CH=CH-CONR^{8c}- or -SO₂NR^{8c}-
 5 wherein R^{8c} is hydrogen atom or C₁₋₆ alkyl.

11. An antagonist according to claim 1, wherein Y is
 an optionally halogenated bivalent C₁₋₆ non-cyclic
 hydrocarbon group.

10

12. An antagonist according to claim 1, wherein Ar is
 a ring of the formula :



15 wherein ----- is a single bond or double bond, n is an
 integer of 1 to 4.

13. (Cancelled)

14. An antagonist according to claim 1, which is an
 20 agent for preventing or treating diseases caused by a
 melanin-concentrating hormone.

15. An antagonist according to claim 1, which is an
 agent for preventing or treating obesity.

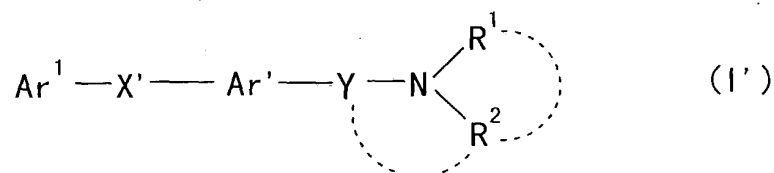
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16. An antagonist according to claim 1, which is an
 anorectic agent.

17. A pharmaceutical composition, which comprises a
 30 melanin-concentrating hormone antagonist in combination
 with at least one species selected from the group
 consisting of an agent for treating diabetes, an agent
 for treating hypertension and an agent for treating

arteriosclerosis and a pharmaceutically acceptable carrier, diluent or excipient.

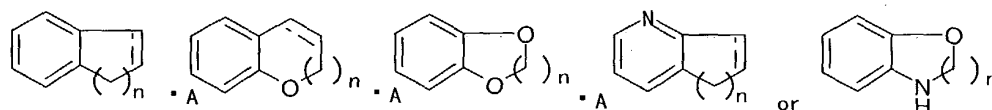
18. A compound of the formula :



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wherein Ar^1 is a cyclic group which may have substituents;

Ar' is a ring of the formula :

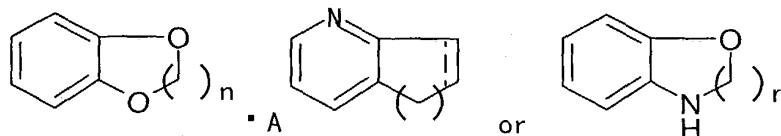


- 10 wherein ----- is a single bond or double bond, n is an integer of 1 to 4, and each ring may have substituents; X' is $-\text{CONR}^{8c}-$, $-\text{NR}^{8c}\text{CO}-$, $-\text{CH}=\text{CH}-\text{CONR}^{8c}-$ or $-\text{SO}_2\text{NR}^{8c}-$

where R^{8c} is hydrogen atom or C_{1-6} alkyl;

Y is a spacer having a main chain of 1 to 6 atoms;

- 15 R^1 and R^2 are independently hydrogen atom or a hydrocarbon group which may have substituents; R^1 and R^2 , together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; or R^2 , together with the adjacent nitrogen atom and Y , may form a nitrogen-containing hetero ring which may have substituents;
- 20 provided that Ar' is a ring of the formula :

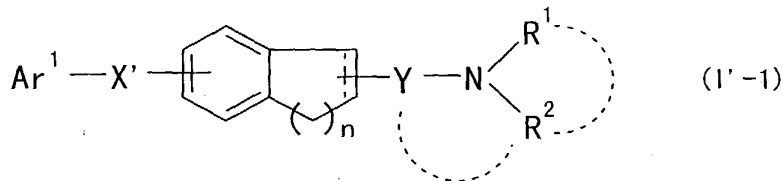


wherein symbols have the same meanings as defined

- 25 above, and each ring may have substituents, when X' is $-\text{SO}_2\text{NH}-$; and provided that Ar^1 is not biphenyl which may be substituted, when X' is $-\text{CONH}-$ and Ar' is any one of benzopyran, dihydrobenzopyran,

dihydrobenzoxazine, dihydrobenzoxazole or tetrahydrobenzoxazepine;
(excluding N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]-4-biphenylcarboxamide); or a salt thereof.

19. A compound of the formula :



wherein Ar¹ is a cyclic group which may have substituents;

----- is a single bond or double bond;

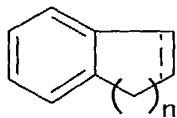
n is an integer of 1 to 4;

X' is -CONR^{8c}-, -NR^{8c}CO- or -CH=CH-CONR^{8c}- where R^{8c} is hydrogen atom or C₁₋₆ alkyl;

Y is a spacer having a main chain of 1 to 6 atoms;

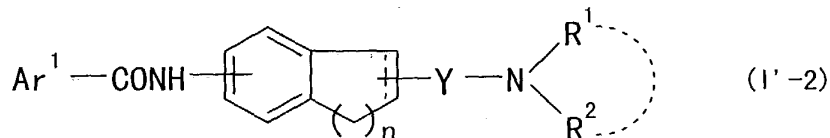
R¹ and R² are independently hydrogen atom or a hydrocarbon group which may have substituents; R¹ and R², together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; or R², together with the adjacent nitrogen atom and Y, may form a nitrogen-containing hetero ring which may have substituents;

a ring of the formula :



wherein symbols have the same meanings as defined above, may have further substituents; provided that N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]-4-biphenylcarboxamide is excluded; or a salt thereof.

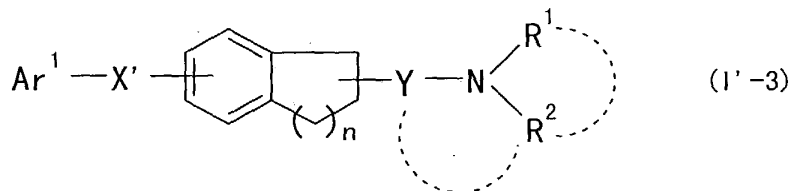
20. A compound according to claim 19, which is of the formula :



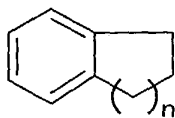
5 wherein R^1 and R^2 are independently hydrogen atom or a hydrocarbon group which may have substituents; R^1 and R^2 , together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; the other symbols have the same meanings
10 as defined in claim 19.

21. (Cancelled)

22. A compound of the formula :

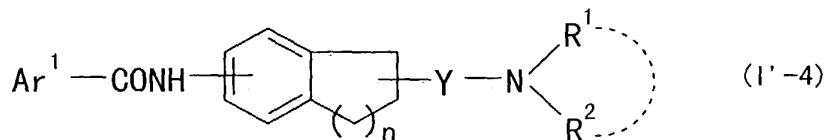


15 wherein Ar^1 is a cyclic group which may have substituents;
n is an integer of 1 to 4;
 X' is $-\text{CONR}^{8c}-$, $-\text{NR}^{8c}\text{CO}-$ or $-\text{CH}=\text{CH}-\text{CONR}^{8c}-$ where R^{8c} is
20 hydrogen atom or C_{1-6} alkyl;
Y is a spacer having a main chain of 1 to 6 atoms;
 R^1 and R^2 are independently hydrogen atom or a hydrocarbon group which may have substituents; R^1 and R^2 , together with the adjacent nitrogen atom, may form
25 a nitrogen-containing hetero ring which may have substituents; or R^2 , together with the adjacent nitrogen atom and Y, may form a nitrogen-containing hetero ring which may have substituents;
a ring of the formula :



- wherein n has the same meaning as defined above, may have further substituents;
provided that N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]-4-biphenylcarboxamide is excluded; or a salt thereof.

23. A compound according to claim 22, which is of the formula :

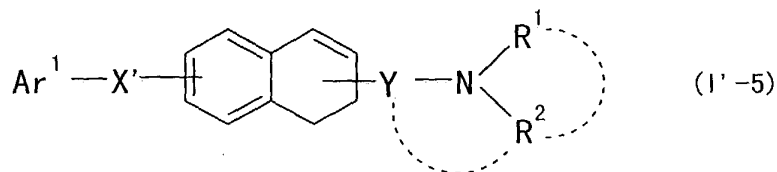


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- wherein R¹ and R² are independently hydrogen atom or a hydrocarbon group which may have substituents; R¹ and R², together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; the other symbols have the same meanings as defined in claim 22.

24. (Cancelled)

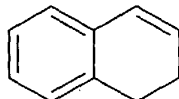
20 25. A compound of the formula :



- wherein Ar¹ is a cyclic group which may have substituents;
X' is -CONR^{8c}-, -NR^{8c}CO- or -CH=CH-CONR^{8c}- where R^{8c} is hydrogen atom or C₁₋₆ alkyl;
Y is a spacer having a main chain of 1 to 6 atoms;
R¹ and R² are independently hydrogen atom or a

25

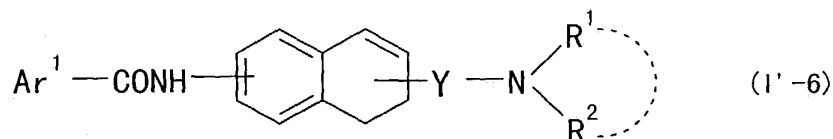
- hydrocarbon group which may have substituents; R^1 and R^2 , together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; or R^2 , together with the adjacent
- 5 nitrogen atom and Y, may form a nitrogen-containing hetero ring which may have substituents;
a ring of the formula :



may have further substituents; or a salt thereof.

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26. A compound according to claim 25, which is of the formula :

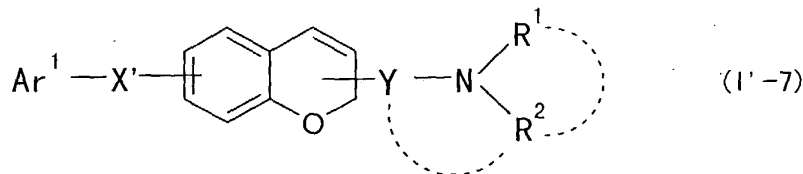


- wherein R^1 and R^2 are independently hydrogen atom or a
- 15 hydrocarbon group which may have substituents; R^1 and R^2 , together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; the other symbols have the same meanings as defined in claim 25.

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27. (Cancelled)

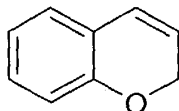
28. A compound of the formula :



- 25 wherein Ar^1 is a cyclic group which may have substituents;

X' is $-CONR^{8c}-$, $-NR^{8c}CO-$, $-CH=CH-CONR^{8c}-$ or $-SO_2NR^{8c}-$
where R^{8c} is hydrogen atom or C_{1-6} alkyl;

Y is a spacer having a main chain of 1 to 6 atoms;
 R^1 and R^2 are independently hydrogen atom or a
hydrocarbon group which may have substituents; R^1 and
 R^2 , together with the adjacent nitrogen atom, may form
5 a nitrogen-containing hetero ring which may have
substituents; or R^2 , together with the adjacent
nitrogen atom and Y, may form a nitrogen-containing
hetero ring which may have substituents;
a ring of the formula :



10

may have further substituents;
provided that Ar^1 is not biphenyl which may be
substituted, when X' is $-CONH-$; or a salt thereof.

15 29. (Cancelled)

30. (Cancelled)

31. (Cancelled)

32. (Cancelled)

33. (Cancelled)

20

34. A compound according to claim 18, which is
N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]-(4'-
methoxybiphenyl-4-yl)carboxamide;

4'-fluoro-N-[6-[(N,N-dimethylamino)methyl]-7,8-dihydro-
25 2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;

4'-fluoro-N-[6-(1-piperidinylmethyl)-7,8-dihydro-2-
naphthalenyl][1,1'-biphenyl]-4-carboxamide;

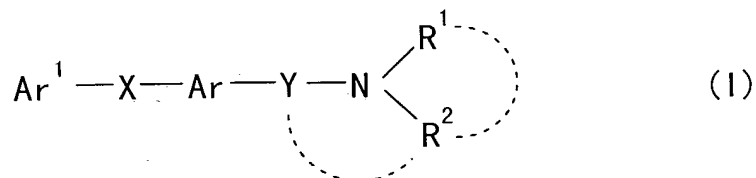
4'-fluoro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-
tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-
30 carboxamide;

(+)-4'-fluoro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-
tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-
carboxamide;

- (-)-4'-fluoro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;
- 4'-chloro-N-[3-[(N,N-dimethylamino)methyl]-2H-chromen-7-yl][1,1'-biphenyl]-4-carboxamide;
- 5 4'-fluoro-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;
- N-[3-[(dimethylamino)methyl]-2H-chromen-7-yl]-4'-fluoro[1,1'-biphenyl]-4-carboxamide;
- 10 4'-chloro-N-[6-[(dimethylamino)methyl]-5-methyl-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;
- 6-(4-methoxyphenyl)-N-[5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]nicotinamide;
- 15 4'-chloro-N-[7-[(dimethylamino)methyl]-5,6-dihydro-3-quinolinyl][1,1'-biphenyl]-4-carboxamide;
- 4-(4-chlorophenyl)-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]-3,6-dihydro-1(2H)-pyridinecarboxamide;
- 20 N-[6-[(dimethylamino)methyl]-7,8-dihydro-2-naphthalenyl]-4-(4-fluorophenyl)-1-piperidinecarboxamide;
- 4-(4-methoxyphenyl)-N-[6-(1-pyrrolidinylmethyl)-5-methyl-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide;
- 25 4'-fluoro-N-[6-[2-(1-pyrrolidinyl)ethyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;
- 4'-chloro-N-[6-[2-(1-pyrrolidinyl)ethyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;
- 30 4'-chloro-N-[2-[(dimethylamino)methyl]-3,4-dihydro-2H-1,4-benzoxazin-6-yl][1,1'-biphenyl]-4-carboxamide;
- 4-(4-methoxyphenyl)-N-[5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide;
- 35 4-(4-chlorophenyl)-N-[6-[(4-methyl-1-

- piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide;
 4'-chloro-N-[2-[(dimethylamino)methyl]-1H-inden-6-yl][1,1'-biphenyl]-4-carboxamide;
 5 4'-fluoro-N-[2-(1-pyrrolidinylmethyl)-3,4-dihydro-2H-1,4-benzoxazin-6-yl][1,1'-biphenyl]-4-carboxamide;
 4'-fluoro-N-[5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;
 10 4'-chloro-N-[5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide; or
 4-(4-chlorophenyl)-N-[5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide.
 15

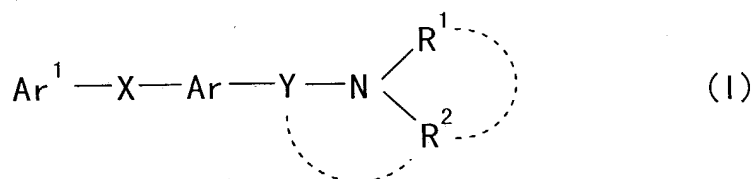
35. A method for preventing or treating diseases caused by a melanin-concentrating hormone in a mammal in need thereof, which comprises administering to said
 20 mammal an effective amount of a compound of the formula
 :



- wherein Ar^1 is a cyclic group which may have substituents;
 25 X is a spacer having a main chain of 1 to 6 atoms;
 Y is a bond or a spacer having a main chain of 1 to 6 atoms;
 Ar is a monocyclic aromatic ring which may be condensed with a 4 to 8 membered non-aromatic ring, and may have
 30 further substituents;
 R^1 and R^2 are independently hydrogen atom or a hydrocarbon group which may have substituents; R^1 and

R², together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; R² may form a spiro ring together with Ar; or R², together with the adjacent nitrogen atom and Y, may form a nitrogen-containing hetero ring which may have substituents; or a salt thereof.

36. A method for preventing or treating obesity in a mammal in need thereof, which comprises administering to said mammal an effective amount of a compound of the formula :



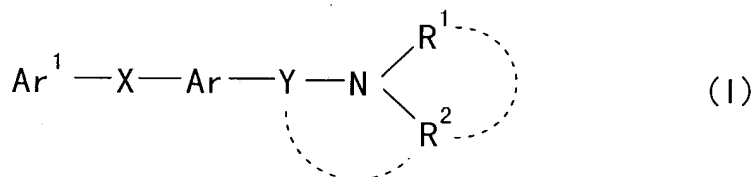
wherein Ar¹ is a cyclic group which may have substituents;

X is a spacer having a main chain of 1 to 6 atoms;
Y is a bond or a spacer having a main chain of 1 to 6 atoms;
Ar is a monocyclic aromatic ring which may be condensed with a 4 to 8 membered non-aromatic ring, and may have further substituents;
R¹ and R² are independently hydrogen atom or a hydrocarbon group which may have substituents; R¹ and R², together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; R² may form a spiro ring together with Ar; or R², together with the adjacent nitrogen atom and Y, may form a nitrogen-containing hetero ring which may have substituents; or a salt thereof.

37. (Cancelled)

38. (Cancelled)

39. A method for antagonizing melanin-concentrating hormone in a mammal in need thereof, comprising administering a compound of the formula :



- 5 wherein Ar¹ is a cyclic group which may have substituents;
 X is a spacer having a main chain of 1 to 6 atoms;
 Y is a bond or a spacer having a main chain of 1 to 6
 10 atoms;
 Ar is a monocyclic aromatic ring which may be condensed with a 4 to 8 membered non-aromatic ring, and may have further substituents;
 R¹ and R² are independently hydrogen atom or a
 15 hydrocarbon group which may have substituents; R¹ and R², together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; R² may form a spiro ring together with Ar; or R², together with the adjacent nitrogen atom and
 20 Y, may form a nitrogen-containing hetero ring which may have substituents; or a salt thereof
 to a mammal.

25 40. A pharmaceutical composition which comprises a compound as defined in claim 18 and a pharmaceutically acceptable carrier, diluent or excipient.

30 41. A pharmaceutical composition which comprises a compound as defined in claim 19, and a pharmaceutically acceptable carrier, diluent or excipient.

42. A pharmaceutical composition which comprises a compound as defined in claim 22 and a pharmaceutically acceptable carrier, diluent or excipient.
- 5 43. A pharmaceutical composition which comprises a compound as defined in claim 25 and a pharmaceutically acceptable carrier, diluent or excipient.
- 10 44. A pharmaceutical composition which comprises a compound as defined in claim 26 and a pharmaceutically acceptable carrier, diluent or excipient.
- 15 45. A pharmaceutical composition which comprises a compound as defined in claim 28 and a pharmaceutically acceptable carrier, diluent or excipient.

DESCRIPTION

Melanin Concentrating Hormone Antagonist

5 TECHNICAL FIELD

The present invention relates to a melanin-concentrating hormone antagonist which is useful as an agent for preventing or treating obesity, etc.

10 BACKGROUND ART

Feeding behavior is an essential action for many living beings including humans. Therefore, if irregularities in feeding behavior occur, disorders, often connected to diseases, will occur in normal life-maintaining activities. Accompanying recent changes of our dietary environment, obesity is now becoming a social problem. In addition, not only is obesity a serious risk factor for life-style diseases such as diabetes, hypertension, and arteriosclerosis; it is also widely known that increased body weight places excessive burdens on joints such as knee joints, causing arthritis and pain.

The "diet boom," etc. show that there is a potentially great percentage of the population hoping to reduce body weight; on the other hand, many cases of feeding problems such as overeating, occurring due to causes such as hereditary neurosis or neurosis due to stress, have been reported.

Therefore, research on and development of agents for preventing or treating obesity, or agents for inhibiting eating, have been vigorously done for a long time.

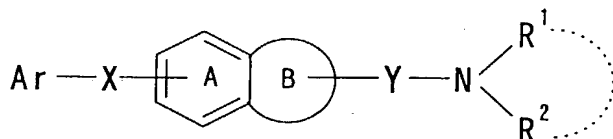
The centrally acting anorectic drug, Mazindol, is now being marketed.

Many appetite control factors such as leptin, have recently been discovered, and the development of anti-obesity agents or anorectic agents which will regulate the functions of these appetite control factors is progressing.

In particular, it is known that melanin- concentrating hormone (hereinafter also abbreviated as "MCH") originates in the hypothalamus and has orexigenic action. In addition, it has been reported that even though the daily behavior of MCH knock-out mice was normal, the amount of feeding by MCH knock-out mice was significantly reduced and their body weights were lighter than those of normal mice [Nature, Vol. 396, p.670, 1998]. This indicates that, if a MCH antagonist was produced, it can be expected to be an excellent anorectic agent or anti-obesity agent; but at present there are no known compound, especially non-peptide type compounds, which possess MCH antagonistic actions.

On the other hand, the following compounds are known as amine derivatives.

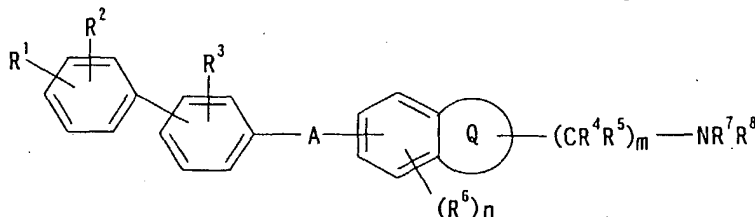
1) W098/38156 describes a compound of the formula :



wherein Ar is an optionally substituted ring assembly aromatic group or an optionally substituted condensed aromatic group; X is a bond, etc.; Y is an optionally substituted bivalent C₁₋₆ aliphatic hydrocarbon group which may have an intervening oxygen atom or sulfur atom; R¹ and R² are independently hydrogen atom or a lower alkyl, or R¹ and R², together with the adjacent nitrogen atom, form an optionally substituted nitrogen-containing hetero ring; Ring A is a benzene ring which may have further substituents in addition to the groups of the formula : -X-Ar where each symbol has the same meaning as defined above; Ring B is a 4 to 8 membered ring which may have further substituents in addition to the group of the formula : -Y-NR¹R² where each symbol has the same meaning as defined above; with the proviso that the condensed ring formed by ring A and ring B is an indole ring, the group of the formula : -X-Ar where

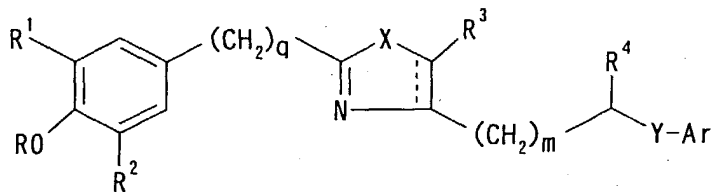
each symbol has the same meaning as defined above is substituted at the 4-, 6-, or 7- position on the indole ring; or its salt, which has an action of inhibiting the production and secretion of β -amyloid protein.

5 2) W095/32967 describes compound of the formula :



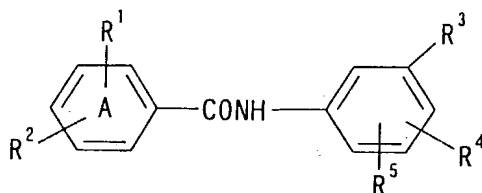
wherein A is CONR, in which R is hydrogen or C₁₋₆ alkyl; Q is an optionally substituted 5 to 7 membered hetero ring containing 1 to 3 hetero atoms selected from nitrogen or sulfur; R¹ is hydrogen, halogen, etc.; R² and R³ are independently hydrogen, halogen, etc.; R₄ and R₅ are independently hydrogen or C₁₋₆ alkyl; R⁶ is halogen, hydroxy, etc.; R₇ and R₈ are independently hydrogen, C₁₋₆ alkyls, etc.; m is 0 to 4; n is 0, 1 or 2; or its salt, which has 5HT1D antagonist activity and can be expected to ameliorate anorexia.

3) W098/15274 describes a compound of the formula :

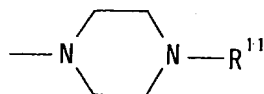


wherein Ar is phenyl, etc.; X is -O- or -S-; Y is CR⁵R^{5'}- where R^{5'} is H and R⁵ is -H, etc.; Z is -CH₂- or -N-; R is H or -(C1-C6) alkyl; R¹ and R² are independently -(C1-C6) alkyl, etc.; R³ is H etc.; R⁴ is hydrogen, etc.; m is an integer of 0 to 2; q is 0 or 1; n is an integer of 0 to 4; p is an integer of 1 to 6; t is an integer of 1 to 4; which has an anti-oxidant activity and can be expected to ameliorate Alzheimer's disease.

4) EP533266



wherein R^1 is halogen, etc.; R^2 is phenyl optionally substituted by 1 or 2 substituents selected from halogen, etc.; R^3 is



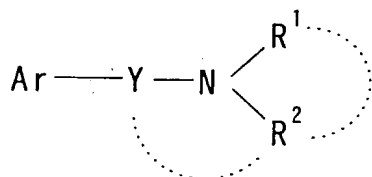
5

; R^4 and R^5 are independently hydrogen, halogen, etc.; R^{11} is hydrogen or C_{1-6} alkyl; which has 5HT_{1D} antagonist activity, and can be expected to ameliorate anorexia.

There has been great desire for the development of a melanin-concentrating hormone antagonist which is useful as an agent for preventing or treating obesity, excellent in oral absorbency, and safe.

DISCLOSURE OF INVENTION

As a result of intensive studies of compounds with a MCH antagonistic action, the present inventors found that a derivative which is obtained by introducing a group of the formula : Ar^1-X- where each symbol has the same meaning as defined hereafter, into a compound of the formula :



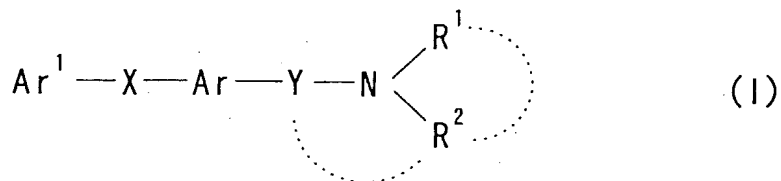
20

wherein each symbol has the same meaning as defined hereinafter, had an excellent MCH antagonistic actions, to complete this invention.

Namely, the present invention relates to :

(1) a melanin-concentrating hormone antagonist which comprises a compound of the formula :

25



wherein Ar¹ is a cyclic group which may have substituents;
X is a spacer having a main chain of 1 to 6 atoms;

Y is a bond or a spacer having a main chain of 1 to 6 atoms;
Ar is a monocyclic aromatic ring which may be condensed with
a 4 to 8 membered non-aromatic ring, and may have further
substituents;

R¹ and R² are independently hydrogen atom or a hydrocarbon
group which may have substituents; R¹ and R², together with
the adjacent nitrogen atom, may form a nitrogen-containing
hetero ring which may have substituents; R² may form a spiro
ring together with Ar; or R², together with the adjacent
nitrogen atom and Y, may form a nitrogen-containing hetero
ring which may have substituents; or a salt thereof;

(2) an antagonist according to the above (1), wherein Y is
a spacer having a main chain of 1 to 6 atoms; R¹ and R² are
independently hydrogen atom or a hydrocarbon group which
may have substituents; R¹ and R², together with the adjacent
nitrogen atom, may form a nitrogen-containing hetero ring
which may have substituents; or R² may form a spiro ring
together with Ar;

(3) an antagonist according to the above (2), wherein Ar¹
is an aromatic group which may have substituents; and "a
hydrocarbon group which may have substituents" for R¹ and
R² is "C₁₋₆ alkyl which may have substituents";

(4) an antagonist according to the above (1), wherein the
cyclic group for Ar¹ is C₆₋₁₄ monocyclic or condensed
polycyclic aromatic hydrocarbon group;

(5) an antagonist according to the above (1), wherein the
cyclic group for Ar¹ is a group formed by removing an
optional one hydrogen atom from an aromatic ring assemble
in which 2 or 3 C₆₋₁₄ monocyclic or condensed polycyclic
aromatic hydrocarbon groups are directly bonded by single

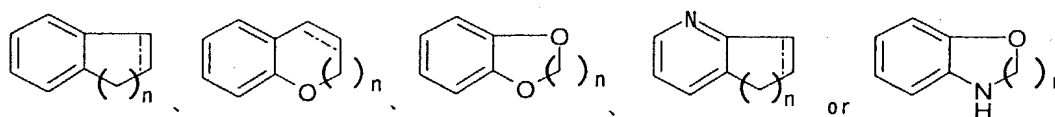
bonds;

(6) an antagonist according to the above (1), wherein the cyclic group for Ar¹ is a group formed by removing an optional one hydrogen atom from an aromatic ring assemble
5 in which C₆₋₁₄ monocyclic or condensed polycyclic aromatic hydrocarbon and 5 to 10 membered aromatic hetero ring are directly bonded by a single bond;

(7) an antagonist according to the above (1), wherein Ar¹ is phenyl, biphenyl, phenyl-pyridyl, phenyl-furyl,
10 phenyl-isoxazolyl, diphenyl-oxazolyl, pyridyl-phenyl, phenyl-pyrimidinyl, benzofuranyl-phenyl, furyl-phenyl, terphenyl, thienyl-phenyl, indolyl, naphthyl-oxadiazolyl, benzofuranyl-oxadiazolyl, benzothienyl, benzofuranyl, fluorenyl, pyridyl-pyrrolyl or
15 thioxanthanyl;

each of which may have 1 to 3 substituents selected from the group consisting of halogen atom; nitro; C₁₋₃ alkylenedioxy; optionally halogenated C₁₋₆ alkyl; hydroxy-C₁₋₆ alkyl; optionally halogenated C₃₋₆ cycloalkyl;
20 optionally halogenated C₁₋₆ alkoxy; optionally halogenated C₁₋₆ alkythio; hydroxy; C₇₋₁₉ aralkyloxy which may have substituents; C₆₋₁₄ aryloxy which may have substituents; amino; mono-C₁₋₆ alkylamino; di-C₁₋₆ alkylamino; 5 to 7 membered saturated cyclic amino which may have substituents
25 and may be condensed with a benzene ring; 5 to 7 membered non-aromatic heterocyclic groups which may have substituents; formyl; carboxy; C₆₋₁₄ aryl-carbonyl which may have substituents; C₆₋₁₄ aryl-carbamoyl which may have substituents; aromatic hetero ring-carbamoyl which may
30 have substituents; C₁₋₆ alkoxy-carbonyl; optionally halogenated C₁₋₆ alkyl-carboxamide; C₆₋₁₄ aryl-carboxamide which may have substituents; C₇₋₁₉ aralkyl-carboxamide which may have substituents; aromatic hetero ring-carboxamide which may have substituents; N-(C₆₋₁₄ aryl-carbonyl which
35 may have substituents)-N-C₁₋₆ alkylamino; C₆₋₁₄ arylamino-carbonylamino which may have substituents; C₆₋₁₄

- arylsulfonylamino which may have substituents; C_{6-14} aryl-carbonyloxy which may have substituents; oxo; carboxy- C_{1-6} alkyl; C_{1-6} alkoxy-carbonyl- C_{1-6} alkyl; C_{7-19} aralkyl which may have substituents; aromatic hetero ring- C_{1-6} alkoxy; and cyano;
- (8) an antagonist according to the above (1), wherein Ar^1 is piperidinyl, piperazinyl, pyrrolidinyl, dihydropyridyl or tetrahydropyridyl; each of which may have 1 or 2 substituents selected from the group consisting of oxo, C_{6-14} aryl which may have substituents, hydroxy, C_{7-19} aralkyloxy-carbonyl, and C_{7-19} aralkyl;
- (9) an antagonist according to the above (1), wherein the "spacer having a main chain of 1 to 6 atoms" for X and Y is a bivalent group consisting of 1 to 3 species selected from -O-, -S-, -CO-, -SO-, -SO₂-, -NR⁸- (R⁸ is hydrogen atom, optionally halogenated C_{1-6} alkyl, optionally halogenated C_{1-6} alkyl-carbonyl, optionally halogenated C_{1-6} alkylsulfonyl), and a bivalent C_{1-6} non-cyclic hydrocarbon group which may have substituents;
- (10) an antagonist according to the above (1), wherein X is -CONR^{8c}-, -NR^{8c}CO-, -CH=CH-CONR^{8c}- or -SO₂NR^{8c}- wherein R^{8c} is hydrogen atom or C_{1-6} alkyl;
- (11) an antagonist according to the above (1), wherein Y is an optionally halogenated bivalent C_{1-6} non-cyclic hydrocarbon group;
- (12) an antagonist according to the above (1), wherein Ar is a ring of the formula :



- wherein ----- is a single bond or double bond, n is an integer of 1 to 4;
- (13) an antagonist according to the above (1), wherein R¹ and R² are hydrogen atom or C_{1-6} alkyl which may have substituents; or R¹ and R², together with the adjacent nitrogen atom, form a 3 to 8 membered nitrogen-containing

hetero ring;

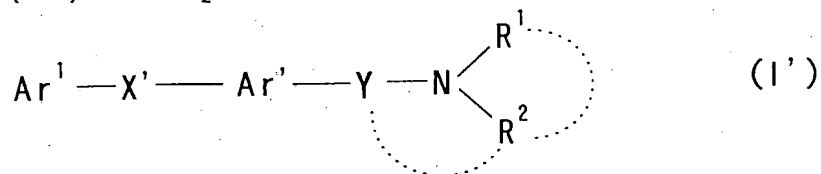
(14) an antagonist according to the above (1), which is an agent for preventing or treating diseases caused by a melanin-concentrating hormone;

5 (15) an antagonist according to the above (1), which is an agent for preventing or treating obesity;

(16) an antagonist according to the above (1), which is an anorectic agent;

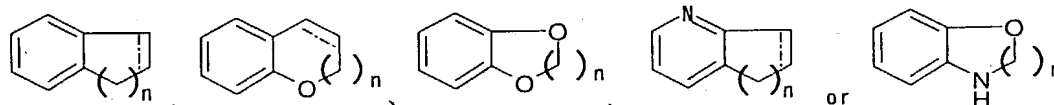
(17) a pharmaceutical, which comprises a melanin-concentrating hormone antagonist in combination with at least one species selected from the group consisting of an agent for treating diabetes, an agent for treating hypertension and an agent for treating arteriosclerosis;

(18) a compound of the formula :



15

wherein Ar^1 is a cyclic group which may have substituents; Ar' is a ring of the formula :



20 wherein ----- is a single bond or double bond, n is an integer of 1 to 4, and each ring may have substituents;

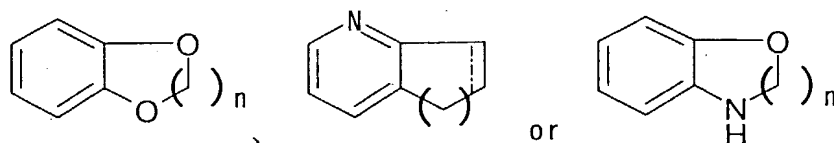
X' is $-\text{CONR}^{\text{sc}}-$, $-\text{NR}^{\text{sc}}\text{CO}-$, $-\text{CH}=\text{CH}-\text{CONR}^{\text{sc}}-$ or $-\text{SO}_2\text{NR}^{\text{sc}}-$ where R^{sc} is hydrogen atom or C_{1-6} alkyl;

Y is a spacer having a main chain of 1 to 6 atoms;

25 R^1 and R^2 are independently hydrogen atom or a hydrocarbon group which may have substituents; R^1 and R^2 , together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; or R^2 , together with the adjacent nitrogen atom and Y , may form a nitrogen-containing hetero ring which may have substituents;

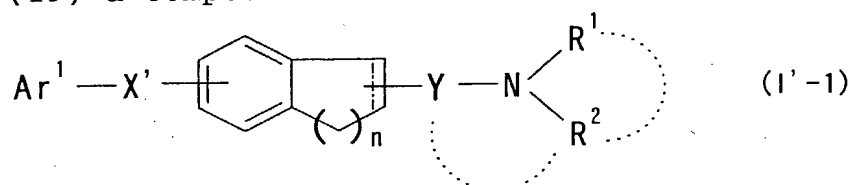
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provided that Ar' is a ring of the formula :



wherein symbols have the same meanings as defined above,
and each ring may have substituents, when X' is -SO₂NH-;
and provided that Ar¹ is not biphenyl which may be
substituted, when X' is -CONH- and Ar' is any one of
benzopyran, dihydrobenzopyran, dihydrobenzoxazine,
dihydrobenzoxazole or tetrahydrobenzoxazepine;
(excluding N-[2-(N,N-dimethylamino)methyl-6-
tetralinyl]-4-biphenylcarboxamide); or a salt thereof;

(19) a compound of the formula :



wherein Ar¹ is a cyclic group which may have substituents;

----- is a single bond or double bond;

n is an integer of 1 to 4;

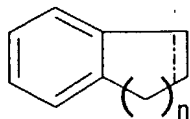
X' is -CONR^{8c}-, -NR^{8c}CO- or -CH=CH-CONR^{8c}- where R^{8c} is
hydrogen atom or C₁₋₆ alkyl;

Y is a spacer having a main chain of 1 to 6 atoms;

R¹ and R² are independently hydrogen atom or a hydrocarbon
group which may have substituents; R¹ and R², together with

the adjacent nitrogen atom, may form a nitrogen-containing
hetero ring which may have substituents; or R², together
with the adjacent nitrogen atom and Y, may form a
nitrogen-containing hetero ring which may have
substituents;

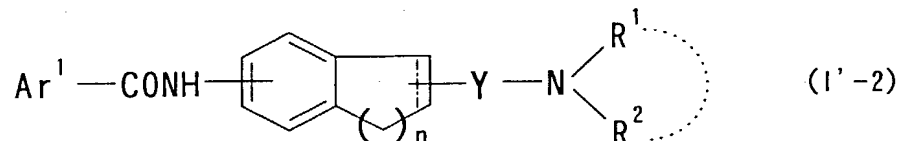
a ring of the formula :



wherein symbols have the same meanings as defined above,

may have further substituents;
provided that N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]-4-biphenylcarboxamide is excluded; or a salt thereof;

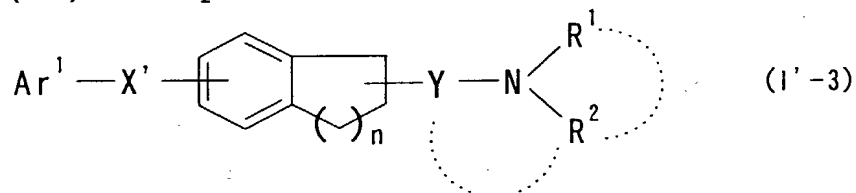
- 5 (20) a compound according to the above (19), which is of the formula :



- wherein R^1 and R^2 are independently hydrogen atom or a hydrocarbon group which may have substituents; R^1 and R^2 , together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; the other symbols have the same meanings as defined in the above (19);

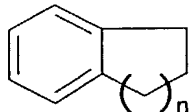
- (21) a compound according to the above (20), wherein Ar^1 is an aromatic group which may have substituents; and "a hydrocarbon group which may have substituents" for R^1 and R^2 is " C_{1-6} alkyl which may have substituents";

(22) a compound of the formula :



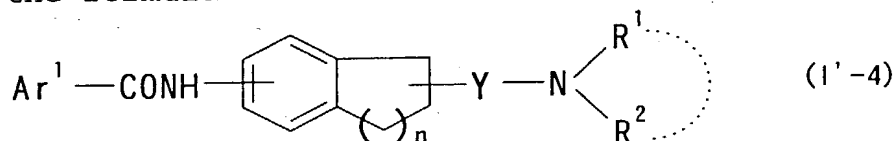
- wherein Ar^1 is a cyclic group which may have substituents;
 n is an integer of 1 to 4;
 X' is $-\text{CONR}^{\text{sc}}-$, $-\text{NR}^{\text{sc}}\text{CO}-$ or $-\text{CH}=\text{CH}-\text{CONR}^{\text{sc}}-$ where R^{sc} is hydrogen atom or C_{1-6} alkyl;
 Y is a spacer having a main chain of 1 to 6 atoms;
25 R^1 and R^2 are independently hydrogen atom or a hydrocarbon group which may have substituents; R^1 and R^2 , together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; or R^2 , together with the adjacent nitrogen atom and Y , may form a
30 nitrogen-containing hetero ring which may have

substituents;
a ring of the formula :



wherein n has the same meaning as defined above, may have
5 further substituents;
provided that N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]-4-biphenylcarboxamide is excluded; or a salt thereof;

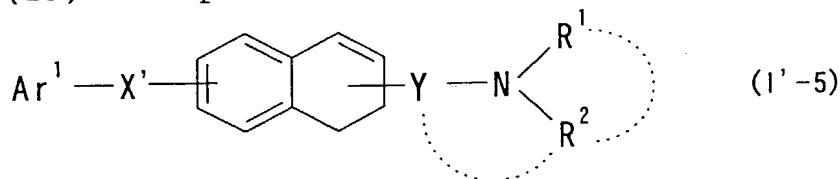
10 (23) a compound according to the above (22), which is of the formula :



wherein R¹ and R² are independently hydrogen atom or a
hydrocarbon group which may have substituents; R¹ and R²,
together with the adjacent nitrogen atom, may form a
15 nitrogen-containing hetero ring which may have substituents; the other symbols have the same meanings as defined in the above (22);

(24) a compound according to the above (23), wherein Ar¹
is an aromatic group which may have substituents; and "a
20 hydrocarbon group which may have substituents" for R¹ and R² is "C₁-₆ alkyl which may have substituents";

(25) a compound of the formula :

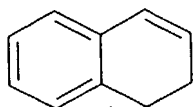


wherein Ar¹ is a cyclic group which may have substituents;
25 X' is -CONR^{8c}-, -NR^{8c}CO- or -CH=CH-CONR^{8c}- where R^{8c} is hydrogen atom or C₁-₆ alkyl;

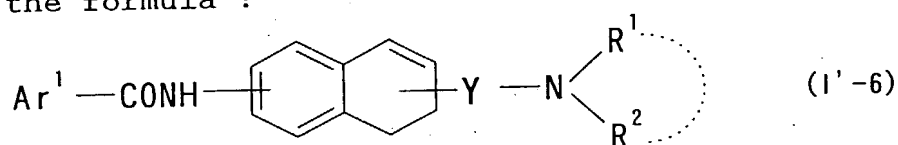
Y is a spacer having a main chain of 1 to 6 atoms;

R¹ and R² are independently hydrogen atom or a hydrocarbon

- group which may have substituents; R^1 and R^2 , together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; or R^2 , together with the adjacent nitrogen atom and Y, may form a
- 5 nitrogen-containing hetero ring which may have substituents;
a ring of the formula :



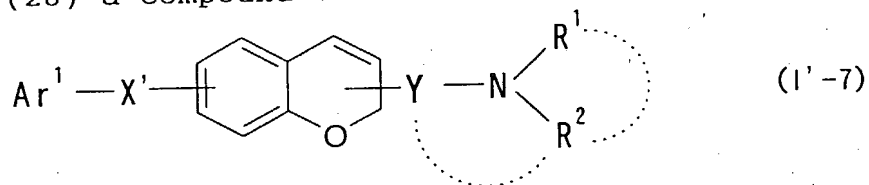
- may have further substituents; or a salt thereof;
- 10 (26) a compound according to the above (25), which is of the formula :



- wherein R^1 and R^2 are independently hydrogen atom or a hydrocarbon group which may have substituents; R^1 and R^2 , together with the adjacent nitrogen atom, may form a
- 15 nitrogen-containing hetero ring which may have substituents; the other symbols have the same meanings as defined in the above (25);

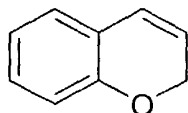
- (27) a compound according to the above (26), wherein Ar^1 is an aromatic group which may have substituents; and "a hydrocarbon group which may have substituents" for R^1 and R^2 is " C_{1-6} alkyl which may have substituents";
- 20

(28) a compound of the formula :

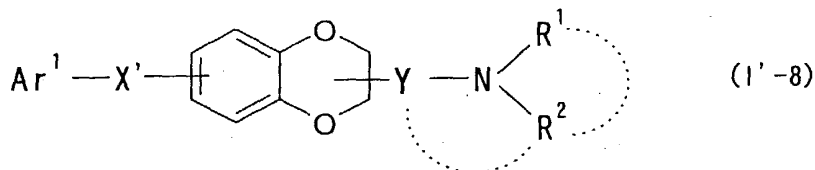


- 25 wherein Ar^1 is a cyclic group which may have substituents;
 X' is $-CONR^{8c}-$, $-NR^{8c}CO-$, $-CH=CH-CONR^{8c}-$ or $-SO_2NR^{8c}-$ where R^{8c} is hydrogen atom or C_{1-6} alkyl;
Y is a spacer having a main chain of 1 to 6 atoms;

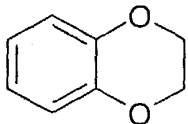
- R^1 and R^2 are independently hydrogen atom or a hydrocarbon group which may have substituents; R^1 and R^2 , together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; or R^2 , together with the adjacent nitrogen atom and Y, may form a nitrogen-containing hetero ring which may have substituents;
- 5 a ring of the formula :



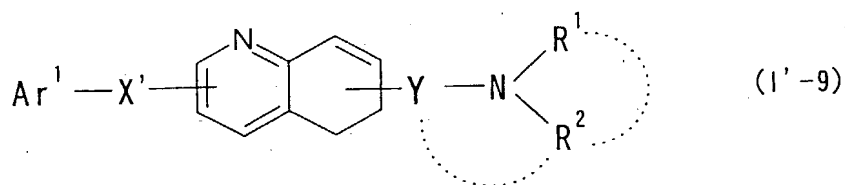
- 10 may have further substituents;
provided that Ar^1 is not biphenyl which may be substituted, when X' is $-CONH-$; or a salt thereof;
(29) a compound of the formula :



- 15 wherein Ar^1 is a cyclic group which may have substituents;
 X' is $-CONR^{8c}-$, $-NR^{8c}CO-$, $-CH=CH-CONR^{8c}-$ or $-SO_2NR^{8c}-$ where R^{8c} is hydrogen atom or C_{1-6} alkyl;
 Y is a spacer having a main chain of 1 to 6 atoms;
 R^1 and R^2 are independently hydrogen atom or a hydrocarbon group which may have substituents; R^1 and R^2 , together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; or R^2 , together with the adjacent nitrogen atom and Y, may form a nitrogen-containing hetero ring which may have substituents;
- 20
25 a ring of the formula :

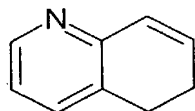


- may have further substituents; or a salt thereof;
(30) a compound of the formula :

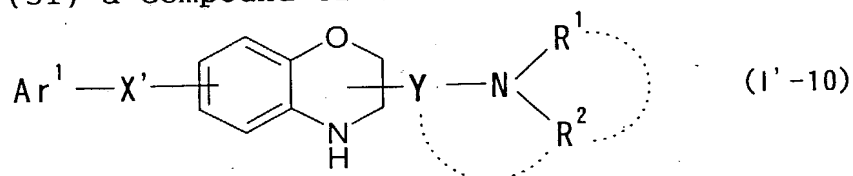


wherein Ar¹ is a cyclic group which may have substituents;
 X' is -CONR^{8c}-, -NR^{8c}CO-, -CH=CH-CONR^{8c}- or -SO₂NR^{8c}- where
 R^{8c} is hydrogen atom or C₁₋₆ alkyl;

- 5 Y is a spacer having a main chain of 1 to 6 atoms;
 R¹ and R² are independently hydrogen atom or a hydrocarbon
 group which may have substituents; R¹ and R², together with
 the adjacent nitrogen atom, may form a nitrogen-containing
 hetero ring which may have substituents; or R², together
 10 with the adjacent nitrogen atom and Y, may form a
 nitrogen-containing hetero ring which may have
 substituents;
 a ring of the formula :



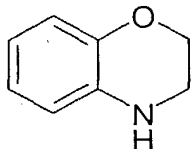
- 15 may have further substituents; or a salt thereof;
 (31) a compound of the formula :



wherein Ar¹ is a cyclic group which may have substituents;
 X' is -CONR^{8c}-, -NR^{8c}CO-, -CH=CH-CONR^{8c}- or -SO₂NR^{8c}- where
 20 R^{8c} is hydrogen atom or C₁₋₆ alkyl;

- Y is a spacer having a main chain of 1 to 6 atoms;
 R¹ and R² are independently hydrogen atom or a hydrocarbon
 group which may have substituents; R¹ and R², together with
 the adjacent nitrogen atom, may form a nitrogen-containing
 hetero ring which may have substituents; or R², together
 25 with the adjacent nitrogen atom and Y, may form a
 nitrogen-containing hetero ring which may have
 substituents;

a ring of the formula :



may have further substituents;

provided that Ar¹ is not biphenyl which may be

5 substituted, when X' is -CONH-; or a salt thereof;

(32) a pharmaceutical composition which comprises a compound as defined in any one of the above (18), (19), (22), (25), (26), (28), (29), (30) and (31);

(33) a prodrug of a compound as defined in any one of the above (18), (19), (22), (25), (26), (28), (29), (30) and (31);

(34) a compound according to the above (18), which is N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]-(4'-methoxybiphenyl-4-yl)carboxamide;

15 4'-fluoro-N-[6-[(N,N-dimethylamino)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;

4'-fluoro-N-[6-(1-piperidinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;

20 4'-fluoro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;

(+)-4'-fluoro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;

25 (-)-4'-fluoro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;

4'-chloro-N-[3-[(N,N-dimethylamino)methyl]-2H-chromen-7-yl][1,1'-biphenyl]-4-carboxamide;

30 4'-fluoro-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;

N-[3-[(dimethylamino)methyl]-2H-chromen-7-yl]-4'-fluoro[1,1'-biphenyl]-4-carboxamide;

- 4'-chloro-N-[6-[(dimethylamino)methyl]-5-methyl-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;
6-(4-methoxyphenyl)-N-[5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]nicotinamide;
5 4'-chloro-N-[7-[(dimethylamino)methyl]-5,6-dihydro-3-quinolinyl][1,1'-biphenyl]-4-carboxamide;
4-(4-chlorophenyl)-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]-3,6-dihydro-1(2H)-pyridinecarboxamide;
10 N-[6-[(dimethylamino)methyl]-7,8-dihydro-2-naphthalenyl]-4-(4-fluorophenyl)-1-piperidinecarboxamide;
4-(4-methoxyphenyl)-N-[6-(1-pyrrolidinylmethyl)-5-methyl-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide;
15 4'-fluoro-N-[6-[2-(1-pyrrolidinyl)ethyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;
4'-chloro-N-[6-[2-(1-pyrrolidinyl)ethyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;
20 4'-chloro-N-[2-[(dimethylamino)methyl]-3,4-dihydro-2H-1,4-benzoxazin-6-yl][1,1'-biphenyl]-4-carboxamide;
4-(4-methoxyphenyl)-N-[5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide;
25 4-(4-chlorophenyl)-N-[6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide;
4'-chloro-N-[2-[(dimethylamino)methyl]-1H-inden-6-yl][1,1'-biphenyl]-4-carboxamide;
30 4'-fluoro-N-[2-(1-pyrrolidinylmethyl)-3,4-dihydro-2H-1,4-benzoxazin-6-yl][1,1'-biphenyl]-4-carboxamide;
4'-fluoro-N-[5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;
35 4'-chloro-N-[5-methyl-6-[(4-methyl-1-

- piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide; or
4-(4-chlorophenyl)-N-[5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide;
- 5 (35) a method for preventing or treating diseases caused by a melanin-concentrating hormone in a mammal in need thereof, which comprises administering to said mammal an effective amount of a compound or a salt thereof as defined in the above (1);
- 10 (36) a method for preventing or treating obesity in a mammal in need thereof, which comprises administering to said mammal an effective amount of a compound or a salt thereof as defined in the above (1);
- 15 (37) use of a compound or a salt thereof as defined in the above (1), for the manufacture of a pharmaceutical preparation for preventing or treating diseases caused by a melanin-concentrating hormone; and
- 20 (38) use of a compound or a salt thereof as defined in the above (1), for the manufacture of a pharmaceutical preparation for preventing or treating obesity.

Examples of "cyclic group" in the "cyclic group which may have substituents" for Ar¹ include aromatic groups, non-aromatic cyclic hydrocarbon groups, non-aromatic heterocyclic groups.

Here, examples of "aromatic groups" include monocyclic aromatic groups, condensed aromatic groups, and ring assembly aromatic groups.

30 Examples of the concerned monocyclic aromatic groups include univalent groups which can be formed by removing an optional one hydrogen atom from a monocyclic aromatic ring. Example of the "monocyclic aromatic ring" include a benzene ring and a 5 or 6 membered aromatic hetero ring.

35 Examples of the "5 or 6 membered aromatic hetero ring" include a 5 or 6 membered aromatic hetero ring containing

one or more (for example, 1 to 3) hetero atom selected from nitrogen, sulfur and oxygen atom in addition to a carbon atom. Concretely, thiophene, furan, pyrrole, imidazole, pyrazole, thiazole, isothiazole, oxazole, isoxazole, pyridine, pyrazine, pyrimidine, pyridazine, 1,2,4-oxadiazole, 1,3,4-oxadiazole, 1,2,4-thiadiazole, 1,3,4-thiadiazole, furazan, etc., can be mentioned.

Concrete examples of the "monocyclic aromatic groups" include phenyl, 2- or 3-thienyl, 2-, 3-, or 4-pyridyl, 2- or 3-furyl, 2-, 4- or 5-thiazonyl, 2-, 4- or 5-oxazolyl, 1-, 3- or 4-pyrazolyl, 2-pyrazinyl, 2-, 4- or 5-pyrimidinyl, 1-, 2- or 3-pyrrolyl, 1-, 2- or 4-imidazolyl, 3- or 4-pyridazinyl, 3-isothiazolyl, 3-isooxazolyl, 1,2,4-oxadiazol-5-yl, 1,2,4-oxadiazol-3-yl.

The "condensed aromatic groups" mean a univalent group that can be formed by removing an optional one hydrogen atom from condensed polycyclic (preferably bicyclic to tetracyclic, more preferably bicyclic or tricyclic) aromatic rings. Examples of the "condensed aromatic groups" include condensed polycyclic aromatic hydrocarbons, condensed polycyclic aromatic hetero rings.

Examples of the "condensed polycyclic aromatic hydrocarbons" include C_{9-14} condensed polycyclic (bicyclic or tricyclic) aromatic hydrocarbons (e.g. naphthalene, indene, fluorene, anthracene, etc.).

Examples of the "condensed polycyclic aromatic hetero rings" include 9 to 14 membered, preferably, 9 or 10 membered, condensed polycyclic aromatic hetero rings containing one or more (for instance, 1 to 4 atoms) hetero atoms selected from nitrogen, sulfur and oxygen atom in addition to carbon atoms. Concrete examples of the "condensed polycyclic aromatic hetero rings" include benzofuran, benzimidazole, benzoxazole, benzothiazole, benzisothiazole, naphtho[2,3-b]thiophene, isoquinoline, quinoline, indole, quinoxaline, phenanthridine, phenothiadine, phenoxazine, phthaladine, naphthylidine,

quinazoline, cinnoline, carbazole, β -carboline, acridine, phenazine, phthalimide, thioxanthene.

Concrete examples of "condensed aromatic groups" include 1-naphthyl; 2-naphthyl; 2-, 3-, 4-, 5- or 8-quinolyl; 1-, 3-, 4-, 5-, 6-, 7- or 8-isoquinolyl; 1-, 2-, 3-, 4-, 5-, 6- or 7-indolyl; 1-, 2-, 4- or 5-isoindolyl; 1-, 5- or 6-phthalazinyl; 2-, 3- or 5-quinoxalynyl; 2-, 3-, 4-, 5- or 6-benzofuranyl; 2-, 4-, 5- or 6-benzothiazolyl; 1-, 2-, 4-, 5- or 6-benzimidazolyl; 1-, 2-, 3- or 4-fluorenyl; thioxanthenyl.

"Ring assembly aromatic group" means a group formed by removing an optional one hydrogen atom from an aromatic ring assemblies in which 2 or more (preferably 2 or 3) aromatic rings are directly bonded by single bonds, and in which the number of bonds which directly bond the rings, is less by one than the number of ring systems.

Examples of the aromatic ring assemblies include an aromatic ring assemblies formed by 2 or 3 (preferably 2) species selected from C_{6-14} monocyclic or condensed polycyclic aromatic hydrocarbons (e.g. benzene and naphthalene) and 5 to 10 membered (preferably 5 or 6 membered) aromatic hetero rings.

Preferable example of the aromatic ring assemblies include aromatic ring assemblies comprising 2 or 3 aromatic rings selected from benzene, naphthalene, pyridine, pyrimidine, thiophene, furan, thiazole, isothiazole, oxazole, isoxazole, 1,2,4-oxadiazole, 1,3,4-oxadiazole, 1,2,4-thiadiazole, 1,3,4-thiadiazole, quinoline, isoquinoline, indole, benzothiophene, benzoxazole, benzothiazole, benzofuran and pyrrole.

Concrete examples of the "ring assembly aromatic groups" include 2-, 3- or 4-biphenyl; 3-(1-naphthyl)-1,2,4-oxadiazol-5-yl; 3-(2-naphthyl)-1, 2, 4-oxadiazol-5-yl; 3-(2-benzofuranyl)-1,2,4-oxadiazol-5-yl; 3-phenyl-1,2,4-oxadiazol-5-yl; 3-(2-benzoxazolyl)-1,2,4-oxadiazol-5-yl; 3-(3-indolyl)-1,2,4-oxadiazol-5-yl; 3-

(2-indolyl)-1,2,4-oxadiazol-5-yl; 4-phenylthiazol-2-yl;
4-(2-benzofuranyl)thiazol-2-yl; 4-phenyl-1,3-oxazol-5-
yl; 5-phenyl-isothiazol-4-yl; 5-phenyloxazol-2-yl; 4-
(2-thienyl)phenyl; 4-(3-thienyl)phenyl; 3-(3-
5 pyridyl)phenyl; 4-(3-pyridyl)phenyl; 6-phenyl-3-pyridyl;
5-phenyl-1,3,4-oxadiazol-2-yl; 4-(2-naphthyl)phenyl; 4-
(2-benzofuranyl)phenyl; 4,4'-terphenyl; 5-phenyl-2-
pyridyl; 2-phenyl-5-pyrimidinyl; 4-(4-pyridyl)phenyl;
2-phenyl-1,3-oxazol-5-yl; 2,4-diphenyl-1,3-oxazol-5-yl;
10 3-phenyl-isoxazol-5-yl; 5-phenyl-2-furyl; 4-(2-
furyl)phenyl; 3-(4-pyridyl)pyrrolyl.

Preferable groups among the above "aromatic groups"
are "C₆₋₁₄ monocyclic or condensed polycyclic aromatic
hydrocarbon groups (preferably, phenyl, etc.)", "a group
15 formed by removing an optional one hydrogen atom from an
aromatic ring assembles in which 2 or 3 C₆₋₁₄ monocyclic or
condensed polycyclic aromatic hydrocarbon groups are
directly bonded by single bonds (preferably, 2-, 3- or
4-biphenyl; 4,4'-terphenyl, etc.)" and "a group formed by
20 removing an optional one hydrogen atom from an aromatic ring
assembles in which a C₆₋₁₄ monocyclic or condensed polycyclic
aromatic hydrocarbon and 5 to 10 membered aromatic hetero
ring are directly bonded by a single bond (preferably,
6-phenyl-3-pyridyl, 5-phenyl-2-pyridyl, etc.)".

25 Examples of "non-aromatic cyclic hydrocarbon groups"
include C₃₋₈ Cycloalkyl, C₃₋₈ cycloalkenyl.

Here, concrete examples of C₃₋₈ cycloalkyl include
cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl,
cycloheptyl, cyclooctyl.

30 Concrete examples of C₃₋₈ cycloalkenyl include
cyclopropenyl, cyclobutenyl, cyclopentenyl,
cyclohexenyl, cycloheptenyl, cyclooctenyl.

Among the above "non-aromatic cyclic hydrocarbon
groups", C₃₋₈ cycloalkyl is preferable, and cyclohexyl is
35 particularly preferable.

Examples of "non-aromatic heterocyclic groups"

include monocyclic non-aromatic heterocyclic groups,
condensed polycyclic non-aromatic heterocyclic groups.

Examples of the "monocyclic non-aromatic heterocyclic groups" include univalent groups formed by removing an optional one hydrogen atom from monocyclic non-aromatic hetero ring. Examples of the "monocyclic non-aromatic heterocyclic groups" include 5 to 8 membered monocyclic non-aromatic heterocyclic groups containing one or more (e.g. 1 to 3) hetero atoms selected from nitrogen, sulfur and oxygen atom in addition to carbon atoms. Concretely, tetrahydrothiophene, tetrahydrofuran, pyrrolidine, imidazoline, imidazolidine, pyrazoline, pyrazolidine, tetrahydrothiazole, tetrahydroisothiazole, tetrahydrooxazole, tetrahydroisoxazole, piperidine, tetrahydropyridine, dihydropyridine, piperazine, morpholine, thiomorpholine, tetrahydropyrimidine, tetrahydropyridazine, hexamethyleneimine, etc. can be mentioned.

"Condensed polycyclic non-aromatic heterocyclic group" means a univalent group formed by removing an optional one hydrogen atom from a condensed polycyclic (preferably bicyclic to tetracyclic, more preferably bicyclic or tricyclic) non-aromatic hetero ring. Examples of the "condensed polycyclic non-aromatic hetero ring" include 9 to 14 membered, preferably 9 or 10 membered condensed polycyclic non-aromatic hetero rings which contain one or more (e.g. 1 to 4) hetero atoms selected from nitrogen, sulfur and oxygen atom in addition to carbon atoms.

Concretely, dihydrobenzofuran, dihydrobenzimidazole, dihydrobenzoxazole, dihydrobenzothiazole, dihydrobenzisothiazole, dihydronaphtho[2,3-b]thiophene, tetrahydroisoquinoline, tetrahydroquinoline, indoline, isoindoline, tetrahydroquinoxaline, tetrahydrophenanthridine, hexahydrophenothiadine, hexahydrophenoxazine,

tetrahydrophthaladine, tetrahydronaphthylidine,
tetrahydroquinazoline, tetrahydrocinnoline,
tetrahydrocarbazole, tetrahydro- β -carboline,
tetrahydroacridine, tetrahydrophenazine,
5 tetrahydrothioxantene, etc., can be mentioned.

Among the above "non-aromatic heterocyclic groups",
"5 to 8 membered monocyclic non-aromatic heterocyclic
groups (preferably piperidinyl; piperazinyl;
pyrrolidinyl; dihydropyridyl; tetrahydropyridyl, etc.)"
10 are preferable.

Examples of "substituents" in the "cyclic group which
may have substituents" for Ar¹ include oxo, halogen atoms
(e.g. fluorine, chlorine, bromine, iodine, etc.), C₁₋₃
15 alkylenedioxy (e.g. methylenedioxy, ethylenedioxy, etc.),
nitro, cyano, optionally halogenated C₁₋₆ alkyl, hydroxy-C₁₋₆
alkyl, carboxy-C₁₋₆ alkyl, C₁₋₆ alkoxy-carbonyl-C₁₋₆ alkyl,
C₆₋₁₄ aryloxy-C₁₋₆ alkyl (e.g. phenoxymethyl, etc.), C₁₋₆
alkyl-C₆₋₁₄ aryl-C₂₋₆ alkenyl (e.g. methylphenylethenyl,
20 etc.), optionally halogenated C₃₋₆ cycloalkyl, optionally
halogenated C₁₋₆ alkoxy, optionally halogenated C₁₋₆
alkylthio, C₇₋₁₉ aralkyl which may have substituents,
hydroxy, C₆₋₁₄ aryloxy which may have substituents, C₇₋₁₉
aralkyloxy which may have substituents, C₆₋₁₄ aryl-carbamoyl
25 which may have substituents, amino, amino-C₁₋₆ alkyl (e.g.
aminomethyl, aminoethyl, aminopropyl, aminobutyl, etc.),
mono-C₁₋₆ alkylamino (e.g. methylamino, ethylamino,
propylamino, isopropylamino, butylamino, etc.), di-C₁₋₆
alkylamino (e.g. dimethylamino, diethylamino,
30 dipropylamino, dibutylamino, ethylmethylamino, etc.),
mono-C₁₋₆ alkylamino-C₁₋₆ alkyl (e.g. methylaminomethyl,
ethylaminomethyl, propylaminomethyl,
isopropylaminoethyl, butylaminoethyl, etc.), di-C₁₋₆
alkylamino-C₁₋₆ alkyl (e.g. dimethylaminomethyl,
35 diethylaminomethyl, dipropylaminomethyl,
diisopropylaminoethyl, dibutylaminoethyl, etc.), 5 to 7

membered saturated cyclic amino which may have substituents, 5 to 7 membered non-aromatic heterocyclic groups which may have substituents, acyl, acylamino, acyloxy, aromatic hetero ring- C_{1-6} alkoxy.

5 The "cyclic group" for Ar^1 may have 1 to 5, preferably 1 to 3, of the above-mentioned substituents at a substitutable position on the cyclic group. When the number of substituents is 2 or more, each substituents can be the same or different.

10 Also, when the "cyclic group" for Ar^1 is a non-aromatic cyclic hydrocarbon group or a non-aromatic heterocyclic group, the "cyclic group" may have as its substituents, C_{6-14} aryl which may have substituents, and 5 to 10 membered aromatic heterocyclic groups which may have substituents.

15 Here, the groups exemplified as "substituents" in the "5 to 7 membered saturated cyclic amino which may have substituents" mentioned hereinafter, can be mentioned as " C_{6-14} aryl which may have substituents" and "5 to 10 membered aromatic heterocyclic groups which may have substituents".

20 The number of substituents is, for instance, 1 to 3. When the number of substituents is 2 or more, each substituents can be the same or different.

Concrete examples of the above "optionally halogenated C_{1-6} alkyl" include C_{1-6} alkyl (e.g. methyl, ethyl, propyl, isopropyl, butyl, isobutyl, sec-butyl, tert-butyl, pentyl, hexyl, etc.) which may have 1 to 5, preferably 1 to 3, halogen atoms (e.g. fluorine, chlorine, bromine, iodine, etc.). Concrete examples include methyl, chloromethyl, difluoromethyl, trichloromethyl, 25 trifluoromethyl, ethyl, 2-bromoethyl, 2,2,2-trifluoroethyl, pentafluoroethyl, propyl, 3,3,3-trifluoropropyl, isopropyl, butyl, 4,4,4-trifluorobutyl, isobutyl, sec-butyl, tert-butyl, pentyl, isopentyl, neopentyl, 5,5,5-trifluoropentyl, hexyl, 6,6,6-trifluorohexyl. 35

The C_{1-6} alkyl in the above "optionally halogenated C_{1-6}

alkyl" can be mentioned as the C_{1-6} alkyl in the above "hydroxy- C_{1-6} alkyl", "carboxy- C_{1-6} alkyl" and " C_{1-6} alkoxy-carbonyl- C_{1-6} alkyl". Examples of C_{1-6} alkoxy in the " C_{1-6} alkoxy-carbonyl- C_{1-6} alkyl" include methoxy, ethoxy, propoxy, butoxy, pentyloxy.

Examples of the above "optionally halogenated C_{3-6} cycloalkyl" include C_{3-6} cycloalkyl (e.g. cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, etc.) which may have 1 to 5, preferably 1 to 3, halogen atoms (e.g. fluorine, chlorine, bromine, iodine, etc.). Concrete examples include cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, 4,4-dichlorocyclohexyl, 2,2,3,3-tetrafluorocyclopentyl, 4-chlorocyclohexyl.

Examples of the above "optionally halogenated C_{1-6} alkoxy" include C_{1-6} alkoxy (e.g. methoxy, ethoxy, propoxy, butoxy, pentyloxy, etc.) which may have 1 to 5, preferably 1 to 3, halogen atoms (e.g. fluorine, chlorine, bromine, iodine, etc.). Concrete examples include methoxy, difluoromethoxy, trifluoromethoxy, ethoxy, 2,2,2-trifluoroethoxy, propoxy, isopropoxy, butoxy, 4,4,4-trifluorobutoxy, isobutoxy, sec-butoxy, pentyloxy, hexyloxy.

Examples of the above "optionally halogenated C_{1-6} alkylthio" include C_{1-6} alkylthio (e.g. methylthio, ethylthio, propylthio, isopropylthio, butylthio, sec-butylthio, tert-butylthio, etc.) which may have 1 to 5, preferably 1 to 3, halogen atoms (e.g. fluorine, chlorine, bromine, iodine, etc.). Concrete examples include methylthio, difluoromethylthio, trifluoromethylthio, ethylthio, propylthio, isopropylthio, butylthio, 4,4,4-trifluorobutylthio, pentylthio, hexylthio.

Examples of the " C_{7-19} aralkyl" in the above " C_{7-19} aralkyl which may have substituents" include benzyl, phenethyl, diphenylmethyl, triphenylmethyl, 1-naphthylmethyl, 2-naphthylmethyl, 2,2-diphenylethyl, 3-phenylpropyl, 4-phenylbutyl, 5-phenylpentyl. Benzyl is

particularly preferable.

Examples of the "substituents" in the above " C_{7-19} aralkyl which may have substituents" include halogen atom (e.g. fluorine, chlorine, bromine, iodine, etc.), C_{1-3} alkylene dioxy (e.g. methylenedioxy, ethylenedioxy, etc.),
5 nitro, cyano, optionally halogenated C_{1-6} alkyl, optionally halogenated C_{3-6} cycloalkyl, optionally halogenated C_{1-6} alkoxy, optionally halogenated C_{1-6} alkylthio, hydroxy, amino, mono- C_{1-6} alkylamino (e.g. methylamino, ethylamino, propylamino, isopropylamino, butylamino, etc.), di- C_{1-6} alkylamino (e.g. dimethylamino, diethylamino, dipropylamino, dibutylamino, ethylmethylamino, etc.),
10 amino- C_{1-6} alkyl (e.g. aminomethyl, aminoethyl, aminopropyl, aminobutyl, etc.), mono- C_{1-6} alkylamino- C_{1-6} alkyl (e.g. methylaminomethyl, ethylaminomethyl, propylaminomethyl, isopropylaminoethyl, butylaminoethyl, etc.), di- C_{1-6} alkylamino- C_{1-6} alkyl (e.g. dimethylaminomethyl, diethylaminomethyl, dipropylaminomethyl, diisopropylaminoethyl, dibutylaminoethyl, etc.), formyl, carboxy, carbamoyl,
20 thiocarbamoyl, optionally halogenated C_{1-6} alkyl-carbonyl, C_{1-6} alkoxy-carbonyl (e.g., methoxycarbonyl, ethoxycarbonyl, propoxycarbonyl, tert-butoxycarbonyl, etc.), mono- C_{1-6} alkyl-carbamoyl (e.g., methylcarbamoyl, ethylcarbamoyl, etc.), di- C_{1-6} alkyl-carbamoyl (e.g. dimethylcarbamoyl, diethylcarbamoyl, ethylmethylcarbamoyl, etc.), optionally halogenated C_{1-6} alkylsulfonyl, formylamino, optionally halogenated C_{1-6} alkyl-carboxamide, C_{1-6} alkoxy-carboxamide (e.g. methoxycarboxamide, ethoxycarboxamide, propoxycarboxamide, butoxycarboxamide, etc.), C_{1-6} alkylsulfonylamino (e.g. methylsulfonylamino, ethylsulfonylamino, etc.), C_{1-6} alkyl-carbonyloxy (e.g. acetoxy, propanoyloxy, etc.), C_{1-6} alkoxy-carbonyloxy (e.g.
35 methoxycarbonyloxy, ethoxycarbonyloxy, propoxycarbonyloxy, butoxycarbonyloxy, etc.) mono- C_{1-6}

alkyl-carbamoyloxy (e.g. methylcarbamoyloxy, ethylcarbamoyloxy, etc.), di- C_{1-6} alkyl-carbamoyloxy (e.g. dimethylcarbamoyloxy, diethylcarbamoyloxy, etc.). The number of substituents is, for instance, 1 to 5, preferably 1 to 3. When the number of substituents is 2 or more, each substituents can be the same or different.

As "optionally halogenated C_{1-6} alkyl", "optionally halogenated C_{3-6} cycloalkyl", "optionally halogenated C_{1-6} alkoxy" and "optionally halogenated C_{1-6} alkylthio", those exemplified as "substituents" in the above "cyclic group which may have substituents" can be used respectively.

Examples of the above "optionally halogenated C_{1-6} alkylcarbonyl" include C_{1-6} alkyl-carbonyl (e.g. acetyl, propanoyl, butanoyl, pentanoyl, hexanoyl, etc.) which may have 1 to 5, preferably 1 to 3, halogen atoms (e.g., fluorine, chlorine, bromine, iodine, etc.). Concrete examples include acetyl, monochloroacetyl, trifluoroacetyl, trichloroacetyl, propanoyl, butanoyl, pentanoyl, hexanoyl.

Examples of the above "optionally halogenated C_{1-6} alkylsulfonyl" include C_{1-6} alkylsulfonyl (e.g. methylsulfonyl, ethylsulfonyl, propylsulfonyl, isopropylsulfonyl, butylsulfonyl, sec-butylsulfonyl, tert-butylsulfonyl, etc.) which may have 1 to 5, preferably 1 to 3, halogen atoms (e.g., fluorine, chlorine, bromine, iodine, etc.). Concrete examples include methylsulfonyl, difluoromethylsulfonyl, trifluoromethylsulfonyl, ethylsulfonyl, propylsulfonyl, isopropylsulfonyl, butylsulfonyl, 4,4,4-trifluorobutylsulfonyl, pentylsulfonyl, hexylsulfonyl.

Examples of the above "optionally halogenated C_{1-6} alkyl-carboxamide" include C_{1-6} alkyl-carboxamide (e.g. acetamide, propanamide, butanamide, etc.) which may have 1 to 5, preferably 1 to 3, halogen atoms (e.g. fluorine, chlorine, bromine, iodine, etc.). Concrete examples include acetamide, trifluoroacetamide, propanamide,

butanamide.

Examples of "C₆₋₁₄ aryloxy" in the above "C₆₋₁₄ aryloxy which may have substituents" include phenyloxy, 1-naphthyloxy, 2-naphthyloxy.

5 Examples of "C₇₋₁₉ aralkyloxy" in the above "C₇₋₁₉ aralkyloxy which may have substituents" include benzyloxy, phenethyloxy, diphenylmethyloxy, triphenylmethyloxy, 1-naphthylmethyloxy, 2-naphthylmethyloxy, 2,2-diphenylethyloxy, 3-phenylpropyloxy, 4-phenylbutyloxy,
10 5-phenylpentyloxy.

Examples of "C₆₋₁₄ arylcarbamoyl" in the above "C₆₋₁₄ arylcarbamoyl which may have substituents" include phenylcarbamoyl, 1-naphthylcarbamoyl, 2-naphthylcarbamoyl.

15 As the "substituents" in the "C₆₋₁₄ aryloxy which may have substituents", "C₇₋₁₉ aralkyloxy which may have substituents" and "C₆₋₁₄ aryl-carbamoyl which may have substituents", those exemplified for "substituents" in the above "C₇₋₁₉ aralkyl which may have substituents" can be used.

20 The number of substituents is, for instance, 1 to 5, preferably 1 to 3. When the number of substituents is 2 or more, each substituents can be the same or different.

Examples of the "5 to 7 membered saturated cyclic amino" in the above "5 to 7 membered saturated cyclic amino
25 which may have substituents" include morpholino, thiomorpholino, piperazin-1-yl, piperidino, pyrrolidin-1-yl. The "5 to 7 membered saturated cyclic amino" can be condensed with a benzene ring.

Examples of "substituents" in the "5 to 7 membered saturated cyclic amino which may have substituents" include
30 oxo, optionally halogenated C₁₋₆ alkyl, optionally halogenated C₁₋₆ alkyl-carbonyl, optionally halogenated C₁₋₆ alkylsulfonyl, C₆₋₁₄ aryl which may have substituents, C₇₋₁₉ aralkyl which may have substituents, C₆₋₁₄ aryl-carbonyl
35 which may have substituents, 5 to 10 membered aromatic heterocyclic group which may have substituents, 5 to 8

membered monocyclic non-aromatic heterocyclic group (e.g., piperidino, piperazinyl, pyrrolidinyl, dihydropyridyl, etc.). The number of substituents is, for instance, 1 to 5, preferably 1 to 3. When the number of substituents is 2 or more, each substituents can be the same or different.

Here, as "optionally halogenated C₁₋₆ alkyl" and "C₇₋₁₉ aralkyl which may have substituents", those exemplified as "substituents" in the above "cyclic group which may have substituents" can be used.

As "optionally halogenated C₁₋₆ alkyl-carbonyl" and "optionally halogenated C₁₋₆ alkylsulfonyl", those exemplified as "substituents" in the above "C₇₋₁₉ aralkyl which may have substituents" can be used.

Examples of the "C₆₋₁₄ aryl" in the "C₆₋₁₄ aryl which may have substituents" include phenyl, 1-naphthyl, 2-naphthyl, 2-indenyl, 2-anthryl. Phenyl is especially preferable.

As the substituents in the "C₆₋₁₄ aryl which may have substituents", those exemplified as "substituents" in the above "C₇₋₁₉ aralkyl which may have substituents" can be used.

The number of substituents is, for instance, 1 to 5, preferably 1 to 3. When the number of substituents is 2 or more, each substituents can be the same or different.

Examples of the "C₆₋₁₄ aryl-carbonyl" in the "C₆₋₁₄ aryl-carbonyl which may have substituents" include benzoyl, 1-naphthoyl, 2-naphthoyl.

As the "substituents" in the "C₆₋₁₄ aryl-carbonyl which may have substituents", those exemplified as "substituents" in the above "C₇₋₁₉ aralkyl which may have substituents" can be used. The number of substituents is, for instance, 1 to 5, preferably 1 to 3. When the number of substituents is 2 or more, each substituents can be the same or different.

Examples of "5 to 10 membered aromatic heterocyclic groups" in "5 to 10 membered aromatic heterocyclic groups which may have substituents" include 5 to 10 membered (monocyclic or bicyclic) aromatic heterocyclic groups

containing 1 or 2 kinds of, preferably 1 to 4 hetero atoms selected from nitrogen, sulfur and oxygen atom in addition to carbon atoms. Concrete examples include 2- or 3-thienyl; 2-, 3- or 4-pyridyl; 2- or 3-furyl; 2-, 4- or 5-thiazolyl; 2-, 4- or 5-oxazolyl; 1-, 3- or 4-pyrazolyl; 2-pyrazinyl; 2-, 4- or 5-pyrimidinyl; 1-, 2- or 3-pyrrolyl; 1-, 2- or 4-imidazolyl; 3- or 4-pyridazinyl; 3-isothiazolyl; 3-isoxazolyl; 1,2,4-oxadiazol-5-yl; 1,2,4-oxadiazol-3-yl; 2-, 3-, 4-, 5- or 8-quinolyl; 1-, 3-, 4-, 5-, 6-, 7- or 8-isoquinolyl; 1-, 2-, 3-, 4-, 5-, 6- or 7-indolyl; 1-, 2-, 4- or 5-isoindolyl; 1-, 5- or 6-phthalazinyl; 2-, 3- or 5-quinoxalinyl; 2-, 3-, 4-, 5- or 6-benzofuranyl; 2-, 4-, 5- or 6-benzothiazolyl; 1-, 2-, 4-, 5- or 6-benzimidazolyl.

Examples of the "substituents" in the "5 to 10 membered aromatic heterocyclic groups which may have substituents" include halogen atom (e.g. fluorine, chlorine, bromine and iodine, etc.), C_{1-3} alkylenedioxy (e.g. methylenedioxy, ethylenedioxy, etc.), nitro, cyano, optionally halogenated C_{1-6} alkyl, C_{6-14} aryloxy- C_{1-6} alkyl (e.g. phenoxymethyl, etc.), C_{1-6} alkyl- C_{6-14} aryl- C_{2-6} alkenyl (e.g. methylphenylethenyl, etc.), optionally halogenated C_{3-6} cycloalkyl, optionally halogenated C_{1-6} alkoxy, optionally halogenated C_{1-6} alkylthio, C_{7-19} aralkyl which may have substituents, hydroxy, C_{6-14} aryloxy which may have substituents, C_{7-19} aralkyloxy which may have substituents, amino, amino- C_{1-6} alkyl (e.g. aminomethyl, aminoethyl, aminopropyl, aminobutyl, etc.), mono- C_{1-6} alkylamino (e.g. methylamino, ethylamino, propylamino, isopropylamino, butylamino, etc.), di- C_{1-6} alkylamino (e.g. dimethylamino, diethylamino, dipropylamino, dibutylamino, ethylmethylamino, etc.), mono- C_{1-6} alkylamino- C_{1-6} alkyl (e.g. methylaminomethyl, ethylaminomethyl, propylaminomethyl, isopropylaminoethyl, butylaminoethyl, etc.), di- C_{1-6} alkylamino- C_{1-6} alkyl (e.g. dimethylaminomethyl, diethylaminomethyl,

dipropylaminomethyl, diisopropylaminoethyl, dibutylaminoethyl, etc.), 5 to 7 membered saturated cyclic amino, acyl, acylamino, acyloxy. The number of substituents is, for instance, 1 to 5, preferably 1 to 3.

5 When the number of substituents is 2 or more, each substituents can be the same or different.

Here, as "optionally halogenated C_{1-6} alkyl", "optionally halogenated C_{3-6} cycloalkyl", "optionally halogenated C_{1-6} alkoxy", "optionally halogenated C_{1-6} alkylthio", " C_{7-19} aralkyl which may have substituents", " C_{6-14} aryloxy which may have substituents", " C_{7-19} aralkyloxy which may have substituents", those exemplified as the "substituent" in the above "cyclic group which may have substituents" can be used respectively.

15 As a "5 to 7 membered saturated cyclic amino", those exemplified as "5 to 7 membered saturated cyclic amino" regarding "5 to 7 membered saturated cyclic amino which may have substituents" which is a "substituent" in the above "5 to 7 membered saturated cyclic amino which may have substituents" can be used.

20 Examples of the above "acyl" include acyl of the formulae : $-CO-R^3$, $-CO-OR^3$, $-CO-NR^3R^4$, $-CS-NR^3R^4$, $-SO_2-R^{3a}$, $-SO-R^{3a}$, $-PO(-OR^3)-OR^4$ or $-PO_2-R^{3a}$ wherein R^3 is (i) hydrogen atom, (ii) a hydrocarbon group which may have substituents, or (iii) a heterocyclic group which may have substituents; R^{3a} is (i) a hydrocarbon group which may have substituents, or (ii) a heterocyclic group which may have substituents; R^4 is hydrogen atom or C_{1-6} alkyl; R^3 and R^{3a} , together with the adjacent nitrogen atom, can form a nitrogen-containing hetero ring which may have substituents.

30 Examples of the "hydrocarbon group" in "hydrocarbon group which may have substituents" for R^3 or R^4 include straight-chain or cyclic hydrocarbon groups (e.g. alkyl, alkenyl, alkynyl, cycloalkyl, aryl, aralkyl, etc.). Among these, C_{1-19} straight chain or cyclic hydrocarbon groups as

shown below are preferable.

a) C_{1-6} alkyl (e.g. methyl, ethyl, propyl, isopropyl, butyl, isobutyl, sec-butyl, tert-butyl, pentyl, hexyl, etc.);

5 b) C_{2-6} alkenyl (e.g., vinyl, allyl, isopropenyl, 2-butenyl, etc.);

c) C_{2-6} alkynyl (e.g. ethynyl, propargyl, 2-butyne, etc.);

d) C_{3-6} cycloalkyl (e.g. cyclopropyl, cyclobutyl, 10 cyclopentyl, cyclohexyl, etc.); the C_{3-6} cycloalkyl can be condensed with one benzene ring;

e) C_{6-14} aryl (e.g. phenyl, 1-naphthyl, 2-naphthyl, 2-indenyl, 2-anthryl, etc.), preferably phenyl;

f) C_{7-19} aralkyl (e.g. benzyl, phenethyl, 15 diphenylmethyl, triphenylmethyl, 1-naphthylmethyl, 2-naphthylmethyl, 2,3-diphenylethyl, 3-phenylpropyl, 4-phenylbutyl, 5-phenylpentyl, etc.), preferably benzyl.

The "hydrocarbon groups" are preferably C_{1-6} alkyl, C_{6-14} aryl, C_{7-19} aralkyl, etc.

20

Examples of the "substituent" in "hydrocarbon groups which may have substituents" include halogen atom (e.g. fluorine, chlorine, bromine, iodine, etc.), C_{1-3} alkylenedioxy (e.g. methylenedioxy, ethylenedioxy, etc.), 25 nitro, cyano, optionally halogenated C_{1-6} alkoxy, optionally halogenated C_{1-6} alkylthio, hydroxy, amino, mono- C_{1-6} alkylamino (e.g. methylamino, ethylamino, propylamino, isopropylamino, butylamino, etc.), di- C_{1-6} alkylamino (e.g. dimethylamino, diethylamino, 30 dipropylamino, dibutylamino, ethylmethylamino, etc.), formyl, carboxy, carbamoyl, thiocarbamoyl, optionally halogenated C_{1-6} alkyl-carbonyl, C_{1-6} alkoxy-carbonyl (e.g., methoxycarbonyl, ethoxycarbonyl, propoxycarbonyl, tert-butoxycarbonyl, etc.), 5 to 10 membered aromatic 35 heterocyclic groups which may have substituents, C_{6-14} aryl-carbonyl which may have substituents, C_{6-14}

aryloxy-carbonyl which may have substituents, C_{7-19}
aralkyloxy-carbonyl which may have substituents, 5 to 6
membered hetero ring-carbonyl which may have substituents,
mono- C_{1-6} alkyl-carbamoyl (e.g. methylcarbamoyl,
5 ethylcarbamoyl, etc.), di- C_{1-6} alkyl-carbamoyl (e.g.
dimethylcarbamoyl, diethylcarbamoyl,
ethylmethylcarbamoyl, etc.), C_{6-14} aryl-carbamoyl which may
have substituents, 5 to 6 membered hetero ring-carbamoyl
which may have substituents, optionally halogenated C_{1-6}
10 alkylsulfonyl, C_{6-14} arylsulfonyl which may have
substituents, formylamino, C_{1-6} alkyl-carbonyloxy (e.g.
acetoxy, propanoyloxy, etc.), C_{6-14} aryl-carbonyloxy which
may have substituents, C_{1-6} alkoxy-carbonyloxy (e.g.
methoxycarbonyloxy, ethoxycarbonyloxy,
15 propoxycarbonyloxy, butoxycarbonyloxy, etc.), mono- C_{1-6}
alkyl-carbamoyloxy (e.g. methylcarbamoyloxy,
ethylcarbamoyloxy, etc.), di- C_{1-6} alkyl-carbamoyloxy (e.g.
dimethylcarbamoyloxy, diethylcarbamoyloxy, etc.), C_{6-14}
aryl-carbamoyloxy which may have substituents,
20 nicotinoyloxy. The number of substituents is, for
instance, 1 to 5, preferably 1 to 3. When the number of
substituents is 2 or more, each substituents can be the same
or different.

Here, as "optionally halogenated C_{1-6} alkoxy",
25 "optionally halogenated C_{1-6} alkylthio" and " C_{6-14} aryl-
carbamoyl which may have substituents", those exemplified
as a "substituent" in the above "cyclic group which may have
substituents" can be used.

As "optionally halogenated C_{1-6} alkyl-carbonyl" and
30 "optionally halogenated C_{1-6} alkylsulfonyl", those
exemplified as a "substituent" in the above " C_{7-19} aralkyl
which may have substituents" can be used.

As the above "5 to 10 membered aromatic heterocyclic
groups which may have substituents" and " C_{6-14} aryl-carbonyl
35 which may have substituents", those exemplified as
"substituent" in the above "5 to 7 membered saturated cyclic

amino which may have substituents" can be used.

Examples of " C_{6-14} aryloxy-carbonyl" in " C_{6-14} aryloxy-carbonyl which may have substituents" include phenyloxycarbonyl, 1-naphthyloxycarbonyl, 2-naphthyloxycarbonyl.

Examples of " C_{7-19} aralkyloxy-carbonyl" in " C_{7-19} aralkyloxy-carbonyl which may have substituents" include benzyloxycarbonyl, phenethyloxycarbonyl, diphenylmethyloxycarbonyl, triphenylmethyloxycarbonyl, 1-naphthylmethyloxycarbonyl, 2-naphthylmethyloxycarbonyl, 2,2-diphenylethyloxycarbonyl, 3-phenylpropyloxycarbonyl, 4-phenylbutyloxycarbonyl, 5-phenylpentyloxycarbonyl.

Examples of "5 to 6 membered hetero ring-carbonyl" in the above "5 to 6 membered hetero ring-carbonyl which may have substituents" include nicotinoyl, isonicotinoyl, 2-thenoyl, 3-thenoyl, 2-furoyl, 3-furoyl, morpholinocarbonyl, piperidinocarbonyl, pyrrolidin-1-ylcarbonyl.

Examples of the "5 to 6 membered hetero ring-carbamoyl" in the above "5 to 6 membered hetero ring-carbamoyl which may have substituents" include morpholinocarbamoyl, piperidinocarbamoyl, 2-pyridylcarbamoyl, 3-pyridylcarbamoyl, 4-pyridylcarbamoyl, 2-thienylcarbamoyl, 3-thienylcarbamoyl.

Examples of " C_{6-14} arylsulfonyl" in the above " C_{6-14} arylsulfonyl which may have substituents" include phenylsulfonyl, 1-naphthylsulfonyl, 2-naphthylsulfonyl.

Examples of " C_{6-14} aryl-carbonyloxy" in the above " C_{6-14} aryl-carbonyloxy which may have substituents" include benzoyloxy, 1-naphthoyloxy, 2-naphthoyloxy.

Examples of " C_{6-14} aryl-carbamoyloxy" in the above " C_{6-14} aryl-carbamoyloxy which may have substituents" include phenylcarbamoyloxy, naphthylcarbamoyloxy.

As the "substituents" in the above " C_{6-14} aryloxy-

carbonyl which may have substituents", " C_{7-19} aralkyloxy-carbonyl which may have substituents", "5 to 6 membered hetero ring-carbonyl which may have substituents", "5 to 6 membered hetero ring-carbamoyl which may have substituents", " C_{6-14} arylsulfonyl which may have substituents", " C_{6-14} aryl-carbonyloxy which may have substituents" and " C_{6-14} aryl-carbamoyloxy which may have substituents", those exemplified as "substituents" in the above " C_{7-19} aralkyl which may have substituents" can be mentioned. The number of the substituents is, for instance, 1 to 5, preferably 1 to 3. When the number of substituents is 2 or more, each substituents can be the same or different.

Examples of "heterocyclic groups" in the "heterocyclic groups which may have substituents" for R^3 or R^{3a} include a 5 to 14 membered (monocyclic, bicyclic or tricyclic) hetero ring containing 1 or 2 kinds of, 1 to 4 hetero atoms selected from nitrogen, sulfur and oxygen atom in addition to carbon atoms. Preferably, univalent groups formed by removing an optional one hydrogen atom from (i) an aromatic hetero ring, (ii) a 5 to 10 membered non-aromatic hetero ring, or (iii) a 7 to 10 membered hetero-bridge ring, can be mentioned.

Here, examples of the "aromatic hetero ring" include a 5 to 14 membered, preferably 5 to 10 membered, aromatic hetero ring containing one or more hetero atom (e.g. 1 to 4) selected from nitrogen, sulfur and oxygen atom in addition to carbon atoms.

Concrete examples include aromatic hetero rings such as thiophene, furan, pyrrole, imidazole, pyrazole, thiazole, isothiazole, oxazole, isoxazole, pyridine, pyrazine, pyrimidine, pyridazine, 1,2,4-oxadiazole, 1,3,4-oxadiazole, 1,2,4-thiadiazole, 1,3,4-thiadiazole, furazan, benzothiophene, benzofuran, benzimidazole, benzoxazole, benzothiazole, benzisothiazole,

naphtho[2,3-b]thiophene, phenoxathiin, indole, isoindole, 1H-indazole, purine, 4H-quinolidine, isoquinoline, quinoline, phthalazine, naphthylidine, quinoxaline, quinazoline, cinnoline, carbazole, β -carboline, phenanthridine, acridine, phenazine, phenothiadine, phenoxazine, phthalimide, etc.; or a ring formed by condensing these rings (preferably monocyclic rings) with one to multiple (preferably 1 or 2) aromatic rings (e.g. benzene ring, etc.).

10 Examples of "5 to 10 membered non-aromatic hetero rings" include 2- or 3-pyrroline, pyrrolidine, 2- or 3-imidazoline, 2-oxazoline, oxazolidine, 2- or 3-pyrazoline, pyrazolidine, 2-thiazoline, piperidine, piperazine, hexamethylenimine, morpholine, thiomorpholine.

15 Examples of "7 to 10 membered hetero-bridge rings" include quinuclidine, 7-azabicyclo[2.2.1]heptane.

 The "hetero cyclic groups" are preferably 5 to 10 membered (monocyclic or bicyclic) heterocyclic groups containing 1 or 2 kinds of, preferably 1 to 4, hetero atoms selected from nitrogen, sulfur and oxygen atom in addition to carbon atoms. Concretely examples include aromatic heterocyclic groups such as 2- or 3-thienyl; 2-, 3- or 4-pyridyl; 2- or 3-furyl; 2-, 4- or 5-thiazolyl; 2-, 4- or 5-oxazolyl; 1-, 3- or 4-pyrazolyl; 2-pyrazinyl; 2-, 4- or 5-pyrimidinyl; 1-, 2- or 3-pyrrolyl; 1-, 2- or 4-imidazolyl; 3- or 4-pyridazinyl; 3-isothiazolyl; 3-isoxazolyl; 1,2,4-oxadiazol-5-yl; 1,2,4-oxadiazol-3-yl; 2-, 3-, 4-, 5- or 8-quinolyl; 1-, 3-, 4-, 5-, 6-, 7- or 8-isoquinolyl; 1-, 2-, 3-, 4-, 5-, 6- or 7-indolyl; 1-, 2-, 4- or 5-isoindolyl; 1-, 5- or 6-phthalazinyl; 2-, 3- or 5-quinoxalinyl; 2-, 3-, 4-, 5- or 6-benzofuranyl; 2-, 3-, 4-, 5- or 6-benzothienyl; 2-, 4-, 5- or 6-benzothiazolyl; 1-, 2-, 4-, 5- or 6-benzimidazolyl; and non-aromatic heterocyclic groups such as 1-, 2- or 3-pyrrolidinyl; 1-, 2-, 4- or 5-imidazolidinyl; 2- or 4-imidazolinyl; 2-, 3- or 4-pyrazolidinyl; piperidino; 2-, 3- or 4-piperidyl; 1-

or 2-piperazinyl; morpholino.

As the "substituents" in the "heterocyclic groups which may have substituents", those exemplified as "substituents" in the above "5 to 10 membered aromatic
5 heterocyclic groups which may have substituents" can be used. The number of substituents is, for instance, 1 to 5, preferably 1 to 3. When the number of substituents is 2 or more, each substituents can be the same or different.

Examples of "C₁₋₆ alkyl" for R⁴ include methyl, ethyl,
10 propyl, isopropyl, butyl, isobutyl, sec-butyl, tert-butyl, pentyl, hexyl.

Examples of "nitrogen-containing hetero ring" in the "nitrogen-containing hetero ring which may have substituents" formed by R³ and R⁴ together with the adjacent
15 nitrogen atoms, include a 5 to 7 membered nitrogen-containing hetero ring which contains at least one nitrogen atom in addition to carbon atoms and may contain 1 to 3 hetero atoms selected from nitrogen, sulfur and oxygen atom. The
20 "nitrogen-containing hetero rings" are preferably piperidine, morpholine, thiomorpholine, piperazine, pyrrolidine, etc.

As the "substituents" in the "nitrogen-containing hetero ring which may have substituents", those exemplified as "substituents" in the above "5 to 10 membered aromatic
25 heterocyclic groups which may have substituents" can be used. The number of substituents is, for instance, 1 to 5, preferably 1 to 3. When the number of substituents is 2 or more, each substituents can be the same or different.

The "acyl" is preferably formyl, carboxy, carbamoyl,
30 optionally halogenated C₁₋₆ alkyl-carbonyl (e.g. acetyl, etc.), C₁₋₆ alkoxy-carbonyl (e.g. methoxycarbonyl, ethoxycarbonyl, propoxycarbonyl, tert-butoxycarbonyl, etc.), C₆₋₁₄ aryl-carbonyl which may have substituents (e.g. benzoyl, 1-naphthoyl, 2-naphthoyl, etc.), C₆₋₁₄ aryloxy-carbonyl which may have substituents (e.g.
35 phenyloxycarbonyl, 1-naphthyloxycarbonyl, 2-

naphthyloxycarbonyl, etc.), C_{7-19} aralkyloxy-carbonyl which may have substituents (e.g. benzyloxycarbonyl, phenethyloxycarbonyl, etc.), a 5 to 6 membered hetero ring-carbonyl which may have substituents (e.g. 5 nicotinoyl, etc.), mono- C_{1-6} alkyl-carbamoyl (e.g. methylcarbamoyl, ethylcarbamoyl, etc.), di- C_{1-6} alkyl-carbamoyl (e.g. dimethylcarbamoyl, diethylcarbamoyl, ethylmethylcarbamoyl, etc.), C_{6-14} aryl-carbamoyl which may have substituents (e.g. phenylcarbamoyl, 4- 10 methoxyphenylcarbamoyl, 3,4-dimethoxyphenylcarbamoyl, etc.), aromatic hetero ring-carbamoyl which may have substituents (e.g. 2-pyridinylcarbamoyl, 2-quinolinylcarbamoyl etc.), optionally halogenated C_{1-6} alkylsulfonyl (e.g. methylsulfonyl, etc.), C_{6-14} 15 arylsulfonyl which may have substituents (e.g. phenylsulfonyl etc.), etc.

Here, as "optionally halogenated C_{1-6} alkyl-carbonyl" and "optionally halogenated C_{7-19} aralkylsulfonyl", those exemplified as "substituents" in the above " C_{7-19} aralkyl 20 which may have substituents" can be used.

As " C_{6-14} aryl-carbonyl which may have substituents", "substituents" in the above "5 to 7 membered saturated cyclic amino which may have substituents" can be used.

As " C_{6-14} aryloxy-carbonyl which may have 25 substituents", " C_{7-19} aralkyloxy-carbonyl which may have substituents", "5 to 6 membered hetero ring-carbonyl which may have substituents", "aromatic hetero ring-carbamoyl which may have substituents" and " C_{6-14} arylsulfonyl which may have substituents", those exemplified as 30 "substituents" in the above "hydrocarbon groups which may have substituents" can be used.

As " C_{6-14} aryl-carbamoyl which may have substituents", those exemplified as "substituents" in the above "cyclic group which may have substituents" can be used. 35

Examples of the above "acylamino" include amino which

is substituted by 1 or 2 of the above "acyl". Preferably, acylamino of the formulae : $-\text{NR}^5-\text{COR}^6$, $-\text{NR}^5-\text{COOR}^{6a}$, $-\text{NR}^5-\text{SO}_2\text{R}^{6a}$, $-\text{NR}^5-\text{CONR}^{6a}\text{R}^{6b}$, $-\text{PO}(-\text{OR}^5)-\text{OR}^6$, or $-\text{PO}_2-\text{R}^6$ wherein R^5 is hydrogen atom or C_{1-6} alkyl; R^6 has the same meaning as the above R^3 ; R^{6a} has the same meaning as the above R^{3a} ; and R^{6b} has the same meaning as R^4], can be mentioned.

As " C_{1-6} alkyl" for R^5 , the same one as in " C_{1-6} alkyl" for the above R^4 can be mentioned.

The "acylamino" is preferably formylamino, optionally halogenated C_{1-6} alkyl-carboxamide (e.g. methylcarboxamide, trifluoromethylcarboxamide, isopropylcarboxamide, etc.), C_{6-14} aryl-carboxamide which may have substituents (e.g. phenylcarboxamide, 2-methoxyphenylcarboxamide, 4-methoxyphenylcarboxamide, etc.), N-(C_{6-14} aryl-carbonyl which may have substituents)-N- C_{1-6} alkylamino (e.g. N-4-methoxybenzoyl-N-methylamino, etc.), C_{7-19} aralkyl-carboxamide which may have substituents (e.g. benzylcarboxamide, etc.), aromatic hetero ring-carboxamide which may have substituents (e.g. benzothiophen-2-ylcarboxamide, etc.), optionally halogenated C_{1-6} alkoxy-carboxamide (e.g. methoxycarboxamide, ethoxycarboxamide, propoxycarboxamide, butoxycarboxamide, etc.), C_{6-14} arylamino-carbonylamino which may have substituents (e.g. phenylaminocarbonylamino, etc.), optionally halogenated C_{1-6} alkylsulfonylamino (e.g. methylsulfonylamino, trifluoromethylsulfonylamino, ethylsulfonylamino, etc.), C_{6-14} arylsulfonylamino which may have substituents (e.g. 4-methoxyphenylsulfonylamino, etc.).

Here, as "substituents" in " C_{6-14} aryl-carboxamide which may have substituents", "N-(C_{6-14} aryl-carbonyl which may have substituents)-N- C_{1-6} arylkylamino", " C_{7-19} aralkyl-carboxamide which may have substituents", "aromatic hetero ring-carboxamide which may have substituents", " C_{6-14} arylamino-carbonylamino which may

have substituents" and "C₆₋₁₄ arylsulfonylamino which may have substituents", those exemplified as "substituents" in the above "C₇₋₁₉ aralkyl which may have substituents" can be mentioned. The number of substituents is, for instance,
5 1 to 5, preferably 1 to 3. When the number of substituents is 2 or more, each substituents can be the same or different.

Examples of the above "acyloxy" include oxy substituted by one of the above "acyl". Preferably,
10 acyloxy of the formulae : -O-COR⁷, -O-COOR⁷, -O-CONHR⁷, -PO(OH)-OR⁷ or -PO₂-R⁷ wherein R⁷ has the same meaning as the above R³, can be mentioned.

The "acyloxy" is preferably optionally halogenated C₁₋₆ alkyl-carbonyloxy (e.g. acetoxy, propanoyloxy, etc.),
15 C₆₋₁₄ aryl-carbonyloxy which may have substituents (e.g. benzoyloxy, 4-methoxybenzoyloxy, etc.), optionally halogenated C₁₋₆ alkoxy-carbonyloxy (e.g. methoxycarbonyloxy, trifluoromethoxycarbonyloxy, ethoxycarbonyloxy, propoxycarbonyloxy,
20 butoxycarbonyloxy, etc.), mono-C₁₋₆ alkyl-carbamoyloxy (e.g. methylcarbamoyloxy, ethylcarbamoyloxy, etc.), di-C₁₋₆ alkyl-carbamoyloxy (e.g. dimethylcarbamoyloxy, diethylcarbamoyloxy, etc.), C₆₋₁₄ aryl-carbamoyloxy which may have substituents (e.g. phenylcarbamoyloxy,
25 naphthylcarbamoyloxy, etc.), nicotinyloxy, etc.

As "substituents" in "C₆₋₁₄ aryl-carbonyloxy which may have substituents" and "C₆₋₁₄ aryl-carbamoyloxy which may have substituents", those exemplified as "substituents" in the above "C₇₋₁₉ aralkyl which may have substituents" can
30 be mentioned. The number of substituents is, for instance, 1 to 5, preferably 1 to 3. When the number of substituents is 2 or more, each substituents can be the same or different.

Examples of the "5 to 7 membered non-aromatic
35 heterocyclic groups which may have substituents", which is "substituents" in "cyclic group which may have

substituents" for Ar¹, include 4,5-dihydro-1,3-oxazol-2-yl, 4,5-dihydro-1,3-thiazol-2-yl, 4,5-dihydro-1H-2-imidazolyl. As "substituents" in the "5 to 7 membered non-aromatic heterocyclic groups which may have substituents", those exemplified as "substituents" in the above "5 to 7 membered saturated cyclic amino which may have substituents" can be used.

As "acyl", "acyloxy" and "acylamino", which are "substituents" in the "cyclic group which may have substituents" for Ar¹, those exemplified as "substituents" in the above "5 to 10 membered aromatic heterocyclic groups which may have substituents" can be used.

Regarding "aromatic hetero ring-C₁₋₆ alkoxy" which is "substituents" in the "cyclic group which may have substituents" for Ar¹, as "aromatic hetero ring", those exemplified as the above R³ can be used. Examples of "C₁₋₆ alkoxy" include methoxy, ethoxy, propoxy, butoxy, pentyloxy.

"Substituents" in the "cyclic group which may have substituents" for Ar¹ are preferably halogen atom (preferably fluorine, chlorine and bromine, etc.); nitro; C₁₋₃ alkylenedioxy (preferably methylenedioxy, etc.); optionally halogenated C₁₋₆ alkyl (preferably, methyl, ethyl, propyl, trifluoromethyl, etc.); hydroxy-C₁₋₆ alkyl (preferably hydroxymethyl, etc.); optionally halogenated C₃₋₆ cycloalkyl (preferably cyclohexyl, etc.); optionally halogenated C₁₋₆ alkoxy (preferably methoxy, ethoxy, etc.); optionally halogenated C₁₋₆ alkylthio (preferably methylthio, etc.); hydroxy; C₇₋₁₉ aralkyloxy which may have substituents (preferably, 1 to 3 substituents selected from halogen atom, optionally halogenated C₁₋₆ alkyl, optionally halogenated C₁₋₆ alkoxy, optionally halogenated C₁₋₆ alkylthio, etc.) (preferably benzyloxy, 4-methoxybenzyloxy, 3-methoxybenzyloxy, 4-fluorobenzyloxy, 4-methylthiobenzyloxy, 4-ethylbenzyloxy, etc.); C₆₋₁₄

aryloxy which may have substituents (preferably, 1 to 3 optionally halogenated C₁₋₆ alkoxy, etc.) (preferably phenyloxy, 4-methoxyphenyloxy, etc.); amino; mono-C₁₋₆ alkylamino (preferably methylamino, etc.); di-C₁₋₆ alkylamino (preferably dimethylamino, etc.); 5 to 7 membered saturated cyclic amino which may have substituents (preferably 1 to 3 oxo) and may be condensed with a benzene ring (preferably 1,3-dioxo-1,3-dihydro-2H-isoindol-2-yl, etc.); 5 to 7 membered non-aromatic heterocyclic groups which may have substituents (preferably 4,5-dihydro-1,3-oxazol-2-yl, etc.); formyl; carboxy; C₆₋₁₄ aryl-carbonyl which may have substituents (preferably benzoyl, etc.); C₆₋₁₄ aryl-carbamoyl which may have substituents (preferably, 1 to 3 optionally halogenated C₁₋₆ alkoxy, etc.) (preferably, phenylcarbamoyl, 4-methoxyphenylcarbamoyl, 3,4-dimethoxyphenylcarbamoyl, etc.); aromatic hetero ring-carbamoyl which may have substituents (preferably 2-pyridinylcarbamoyl, 2-quinolinylcarbamoyl, etc.); C₁₋₆ alkoxy-carbonyl (preferably methoxycarbonyl, ethoxycarbonyl, etc.); optionally halogenated C₁₋₆ alkyl-carboxamide (preferably methylcarboxamide, trifluoromethylcarboxamide, isopropylcarboxamide, etc.); C₆₋₁₄ aryl-carboxamide which may have substituents (preferably, 1 to 3 optionally halogenated C₁₋₆ alkoxy, etc.) (preferably phenylcarboxamide, 2-methoxyphenylcarboxamide, 4-methoxyphenylcarboxamide, etc.); C₇₋₁₉ aralkyl-carboxamide which may have substituents (preferably benzylcarboxamide, etc.); aromatic hetero ring-carboxamide which may have substituents (preferably benzothiophen-2-ylcarboxamide, etc.); N-(C₆₋₁₄ aryl-carbonyl which may have substituents (preferably, 1 to 3 optionally halogenated C₁₋₆ alkoxy, etc.))-N-C₁₋₆ alkylamino (preferably N-4-methoxybenzoyl-N-methylamino, etc.); C₆₋₁₄ arylamino-carbonylamino which may have substituents (preferably phenylaminocarbonylamino, etc.); C₆₋₁₄ arylsulfonylamino which may have substituents (preferably,

1 to 3 optionally halogenated C_{1-6} alkoxy, etc.) (preferably 4-methoxyphenylsulfonylamino, etc.); C_{6-14} aryl-carbonyloxy which may have substituents (preferably, 1 to 3 optionally halogenated C_{1-6} alkoxy, etc.) (preferably 4-methoxybenzoyloxy, etc.); oxo; carboxy- C_{1-6} alkyl (preferably carboxyethyl, etc.); C_{1-6} alkoxy-carbonyl- C_{1-6} alkyl (preferably methoxycarbonylmethyl, etc.); C_{7-19} aralkyl which may have substituents (preferably 1 to 3 halogen atom) (preferably benzyl, 2,4-dichlorobenzyl, etc.); aromatic hetero ring- C_{1-6} alkoxy (preferably 2-quinolylmethoxy, etc.); cyano, etc.

When "cyclic group" in "cyclic group which may have substituents" for Ar^1 is a non-aromatic cyclic hydrocarbon group or a non-aromatic heterocyclic group, C_{6-14} aryl which may have substituents (preferably, 1 to 3 substituents selected from halogen atom, C_{1-3} alkylenedioxy, optionally halogenated C_{1-6} alkyl, optionally halogenated C_{1-6} alkoxy, etc.) (preferably phenyl, 4-fluorophenyl, 1,3-benzodioxol-5-yl, 4-chlorophenyl, 4-methylphenyl, 4-methoxyphenyl), hydroxy, C_{7-19} aralkyloxy-carbonyl (preferably benzyloxycarbonyl), C_{7-19} aralkyl (preferably benzyl), etc., can be used as a preferable substituent.

Ar^1 is preferably phenyl, biphenyl (preferably 4-biphenyl, 2-biphenyl), phenyl-pyridyl (preferably 6-phenyl-3-pyridyl, 5-phenyl-2-pyridyl), phenyl-furyl (preferably 5-phenyl-2-furyl), phenyl-isoxazolyl (preferably 3-phenyl-isoxazol-5-yl), diphenyl-oxazolyl (preferably 2,4-diphenyl-1,3-oxazol-5-yl), pyridyl-phenyl (preferably 4-(4-pyridyl)phenyl, 4-(3-pyridyl)phenyl), phenyl-pyrimidinyl (preferably 2-phenyl-5-pyrimidinyl), benzofuranyl-phenyl (preferably 4-(2-benzofuranyl)phenyl), furyl-phenyl (preferably 4-(2-furyl)phenyl), terphenyl (preferably 4,4'-terphenyl), thienyl-phenyl (preferably 4-(2-thienyl)phenyl), indolyl (preferably 2-indolyl, 3-indolyl), naphthyl-oxadiazolyl

(preferably 3-(2-naphthyl)-1,2,4-oxadiazol-5-yl), benzofuranyl-oxadiazole (preferably 3-(2-benzofuranyl)-1,2,4-oxadiazol-5-yl), benzothienyl (preferably 2-benzothienyl), benzofuranyl (preferably 2-benzofuranyl), fluorenyl (preferably 2-fluorenyl), pyridyl-pyrrolyl (preferably 3-(4-pyridyl)pyrrolyl), thioxanthenyl; each of which may have 1 to 3 (preferably 1 or 2) substituents selected from the group consisting of halogen atom (preferably fluorine, chlorine, bromine, etc.); nitro; C₁₋₃ alkylenedioxy (preferably methylenedioxy, etc.); optionally halogenated C₁₋₆ alkyl (preferably methyl, ethyl, propyl, trifluoromethyl, etc.); hydroxy-C₁₋₆ alkyl (preferably hydroxymethyl, etc.); optionally halogenated C₃₋₆ cycloalkyl (preferably cyclohexyl, etc.); optionally halogenated C₁₋₆ alkoxy (preferably methoxy, ethoxy, etc.); optionally halogenated C₁₋₆ alkythio (preferably methylthio, etc.); hydroxy; C₇₋₁₉ aralkyloxy which may have substituents (preferably, 1 to 3 substituents selected from halogen atom, optionally halogenated C₁₋₆ alkyl, optionally halogenated C₁₋₆ alkoxy, optionally halogenated C₁₋₆ alkylthio, etc.) (preferably benzyloxy, 4-methoxybenzyloxy, 3-methoxybenzyloxy, 4-fluorobenzyloxy, 4-methylthiobenzyloxy, 4-ethylbenzyloxy, etc.); C₆₋₁₄ aryloxy which may have substituents (preferably, 1 to 3 optionally halogenated C₁₋₆ alkoxy, etc.) (preferably phenyloxy, 4-methoxyphenyloxy, etc.); amino; mono-C₁₋₆ alkylamino (preferably methylamino, etc.); di-C₁₋₆ alkylamino (preferably dimethylamino, etc.); 5 to 7 membered saturated cyclic amino which may have substituents (preferably 1 to 3 oxo) and may be condensed with a benzene ring (preferably 1,3-dioxo-1,3-dihydro-2H-isoindol-2-yl, etc.); 5 to 7 membered non-aromatic heterocyclic groups which may have substituents (preferably 4,5-dihydro-1,3-oxazol-2-yl, etc.); formyl; carboxy; C₆₋₁₄ aryl-carbonyl which may have substituents (preferably benzoyl, etc.); C₆₋₁₄ aryl-carbamoyl which may have substituents

- (preferably, 1 to 3 optionally halogenated C_{1-6} alkoxy, etc.) (preferably phenylcarbamoyl, 4-methoxyphenylcarbamoyl, 3,4-dimethoxyphenylcarbamoyl, etc.); aromatic hetero ring-carbamoyl which may have substituents (e.g. 2-
 5 pridinylcarbamoyl, 2-quinolinylcarbamoyl, etc.); C_{1-6} alkoxy-carbonyl (preferably methoxycarbonyl, ethoxycarbonyl, etc.); optionally halogenated C_{1-6} alkyl-carboxamide (preferably, methylcarboxamide, trifluoromethylcarboxamide, isopropylcarboxamide, etc.);
 10 C_{6-14} aryl-carboxamide which may have substituents (preferably, 1 to 3 optionally halogenated C_{1-6} alkoxy, etc.) (preferably phenylcarboxamide, 2-methoxyphenylcarboxamide, 4-methoxyphenylcarboxamide, etc.); C_{7-19} aralkyl-carboxamide which may have substituents
 15 (preferably benzylcarboxamide, etc.); aromatic hetero ring-carboxamide which may have substituents (preferably benzothiophen-2-ylcarboxamide, etc.); N-(C_{6-14} aryl-carbonyl which may have substituents (preferably, 1 to 3 optionally halogenated C_{1-6} alkoxy, etc.))-N- C_{1-6} alkylamino
 20 (preferably N-4-methoxybenzoyl-N-methylamino, etc.); C_{6-14} arylamino-carbonylamino which may have substituents (preferably phenylaminocarbonylamino, etc.); C_{6-14} arylsulfonylamino which may have substituents (preferably, 1 to 3 optionally halogenated C_{1-6} alkoxy, etc.) (preferably
 25 4-methoxyphenylsulfonylamino, etc.); C_{6-14} aryl-carbonyloxy which may have substituents (preferably, 1 to 3 optionally halogenated C_{1-6} alkoxy, etc.) (preferably 4-methoxybenzoyloxy, etc.); oxo; carboxy- C_{1-6} alkyl (preferably carboxyethyl, etc.); C_{1-6} alkoxy-carbonyl- C_{1-6}
 30 alkyl (preferably methoxycarbonylmethyl, etc.); C_{7-19} aralkyl which may have substituents (preferably 1 to 3 halogen atom) (preferably benzyl, 2,4-dichlorobenzyl, etc.); aromatic hetero ring- C_{1-6} alkoxy (preferably 2-quinolylmethoxy, etc.); and cyano.
- 35 Further, preferable examples of Ar^1 include piperidinyl (preferably piperidino), piperazinyl,

pyrrolidinyl, dihydropyridyl, tetrahydropyridyl; each of which may have 1 or 2 substituents selected from the group consisting of oxo, C₆₋₁₄ aryl which may have substituents (preferably, 1 to 3 substituents selected from halogen atom, C₁₋₃ alkylenedioxy, optionally halogenated C₁₋₆ alkyl, 5 optionally halogenated C₁₋₆ alkoxy, etc.) (preferably phenyl, 4-fluorophenyl, 1,3-benzodioxol-5-yl, 4-chlorophenyl, 4-methylphenyl, 4-methoxyphenyl), hydroxy, C₇₋₁₉ aralkyloxy-carbonyl (preferably benzyloxycarbonyl) and C₇₋₁₉ aralkyl (preferably benzyl).

Ar¹ is more preferably, phenyl, biphenyl (preferably 4-biphenyl) or phenyl-pyridyl (preferably 6-phenyl-3-pyridyl, 5-phenyl-2-pyridyl); each of which may have 1 or 2 substituents selected from the group consisting of 15 halogen atom (preferably fluorine, chlorine, bromine, etc.); optionally halogenated C₁₋₆ alkyl (preferably methyl, ethyl, propyl, trifluoromethyl, etc.); optionally halogenated C₁₋₆ alkoxy (preferably methoxy, ethoxy, etc.); C₇₋₁₉ aralkyloxy which may have substituents (preferably, 20 1 to 3 substituents selected from halogen atom, optionally halogenated C₁₋₆ alkyl, optionally halogenated C₁₋₆ alkoxy, optionally halogenated C₁₋₆ alkylthio, etc.) (preferably benzyloxy, 4-methoxybenzyloxy, etc.); C₆₋₁₄ aryloxy which may have substituents (preferably, 1 to 3 optionally 25 halogenated C₁₋₆ alkoxy, etc.) (preferably phenyloxy, etc.); C₆₋₁₄ aryl-carbonyl which may have substituents (preferably, 1 to 3 optionally halogenated C₁₋₆ alkoxy, etc.) (preferably benzoyl, etc.); C₆₋₁₄ aryl-carbamoyl which may have substituents (preferably, 1 to 3 optionally 30 halogenated C₁₋₆ alkoxy, etc.) (preferably phenylcarbamoyl, 4-methoxyphenylcarbamoyl, 3,4-dimethoxyphenylcarbamoyl, etc.); aromatic hetero ring-carbamoyl which may have substituents (e.g. 2-pyridinylcarbamoyl, 2-quinolinylcarbamoyl, etc.); C₆₋₁₄ aryl-carboxamide which 35 may have substituents (preferably, 1 to 3 optionally halogenated C₁₋₆ alkoxy, etc.) (preferably

phenylcarboxamide, 2-methoxyphenylcarboxamide, 4-methoxyphenylcarboxamide, etc.); C_{7-19} aralkyl-carboxamide which may have substituents (preferably benzylcarboxamide, etc.); aromatic hetero ring-carboxamide (preferably benzothiophen-2-ylcarboxamide, etc.); N-(C_{6-14} aryl-carbonyl which may have substituents (preferably, 1 to 3 optionally halogenated C_{1-6} alkoxy, etc.))-N- C_{1-6} alkylamino (preferably N-4-methoxybenzoyl-N-methylamino, etc.); C_{6-14} arylamino-carbonylamino which may have substituents (preferably phenylaminocarbonylamino, etc.); C_{6-14} arylsulfonylamino which may have substituents (preferably, 1 to 3 optionally halogenated C_{1-6} alkoxy, etc.) (preferably 4-methoxyphenylsulfonylamino, etc.); and C_{6-14} arylcarbonyloxy which may have substituents (preferably, 1 to 3 optionally halogenated C_{1-6} alkoxy, etc.) (preferably 4-methoxybenzoyloxy, etc.).

Further, preferable examples of Ar^1 include piperidino, piperazinyl or pyrrolidinyl; each of which may have 1 or 2 substituents selected from the group consisting of oxo and C_{6-14} aryl (preferably phenyl) which may have substituents [preferably halogen atom (preferably fluorine, chlorine, bromine, etc.), optionally halogenated C_{1-6} alkyl (preferably methyl, ethyl, propyl, trifluoromethyl, etc.) or optionally halogenated C_{1-6} alkoxy (preferably methoxy, ethoxy, etc.)].

The "spacer having a main chain of 1 to 6 atoms" means a space in which 1 to 6 atoms are linked. Here, the "number of atoms in the main chain" is counted so that the number of atoms in the main chain is minimum. For instance, the number of atoms of 1,2-cyclopentylene is counted as 2, and the number of atoms of 1,3-cyclopentylene is counted as 3.

Examples of the "spacer having a main chain of 1 to 6 atoms" include a bivalent group consisting of 1 to 3 species selected from -O-, -S-, -CO-, -SO-, -SO₂-, -NR⁸- (R⁸ is hydrogen atom, optionally halogenated C_{1-6} alkyl,

optionally halogenated C_{1-6} alkyl-carbonyl, optionally halogenated C_{1-6} alkylsulfonyl), bivalent C_{1-6} non-cyclic hydrocarbon groups which may have substituents, and bivalent C_{5-8} monocyclic non-aromatic hydrocarbon groups.

5 Here, as "optionally halogenated C_{1-6} alkyl", those exemplified as "substituents" in the above "cyclic group which may have substituents" can be used.

As "optionally halogenated C_{1-6} alkyl-carbonyl" and "optionally halogenated C_{1-6} alkylsulfonyl", those
10 exemplified as "substituents" in the above " C_{7-19} aralkyl which may have substituents" can be used.

Examples of "bivalent C_{1-6} non-cyclic hydrocarbon groups" in the "bivalent C_{1-6} non-cyclic hydrocarbon groups
15 which may have substituents" include

(1) C_{1-6} alkylene (e.g. $-CH_2-$, $-(CH_2)_2-$, $-(CH_2)_3-$, $-(CH_2)_4-$, $-(CH_2)_5-$, $-(CH_2)_6-$, $-CH(CH_3)-$, $-C(CH_3)_2-$, $-(CH(CH_3))_2-$, $-(CH_2)_2C(CH_3)_2-$, $-(CH_2)_3C(CH_3)_2-$, etc.);

(2) C_{2-6} alkenylene (e.g. $-CH=CH-$, $-CH_2-CH=CH-$, $-C(CH_3)_2-CH=CH-$, $-CH_2-CH=CH-CH_2-$, $-CH_2-CH_2-CH=CH-$, $-CH=CH-CH=CH-$, $-CH=CH-CH_2-CH_2-CH_2-$, etc.);

(3) C_{2-6} alkynylene (e.g. $-C\equiv C-$, $-CH_2-C\equiv C-$, $-CH_2-C\equiv C-CH_2-$, etc.)

each of which may have 1 to 5, preferably 1 to 3, halogen
25 atoms (e.g. fluorine, chlorine, bromine, iodine, etc.).

The "bivalent C_{1-6} non-cyclic hydrocarbon groups" may have 1 to 5, preferably 1 to 3 substituents at a substitutable position. Examples of such substituents include halogen atom (e.g. fluorine, chlorine, bromine, iodine, etc.), hydroxy, C_{1-6} alkyl-carbonyloxy (e.g.,
30 acetoxy, etc.).

As the "bivalent C_{5-8} monocyclic non-aromatic hydrocarbon groups", for instance, bivalent groups formed
35 by removing an optional two hydrogen atoms from C_{5-8} cycloalkane or C_{5-8} cycloalkene, can be mentioned. Concrete

examples include 1,2-cyclopentylene; 1,3-cyclopentylene;
 1,2-cyclohexylene; 1,3-cyclohexylene; 1,4-cyclohexylene;
 1,2-cycloheptylene; 1,3-cycloheptylene; 1,4-
 cycloheptylene; 3-cyclohexen-1,4-ylene; 3-cyclohexen-
 5 1,2-ylene; 2,5-cyclohexadien-1,4-ylene. Especially, C₅₋₈
 cycloalkylene is preferable.

The "spacer having a main chain of 1 to 6 atoms" is
 preferably a bivalent group consisting of 1 to 3 species
 selected from -O-, -S-, -CO-, -SO-, -SO₂-, -NR^a- (R^a has
 10 the same meaning as defined above) and optionally
 halogenated bivalent C₁₋₆ non-cyclic hydrocarbon groups.

Preferred examples of the "spacer having a main chain
 of 1 to 6 atoms" include

(1) C₁₋₆ alkylene (e.g. -CH₂-, -(CH₂)₂-, -(CH₂)₃-, -
 15 (CH₂)₄-, -(CH₂)₅-, -(CH₂)₆-, -CHCH₃-, -C(CH₃)₂-, -CH(CF₃)-,
 -(CH(CH₃))₂-, -(CF₂)₂-, -(CH₂)₂C(CH₃)₂-, -(CH₂)₃C(CH₃)₂-,
 etc.);

(2) C₂₋₆ alkenylene (e.g. -CH=CH-, -CH₂-CH=CH-, -
 CH₂-CF=CH-, -C(CH₃)₂-CH=CH-, -CH₂-CH=CH-CH₂-, -CH₂-CH₂-
 20 CH=CH-, -CH=CH-CH=CH-, -CH=CH-CH₂-CH₂-CH₂-, etc.);

(3) C₂₋₆ alkynylene (e.g. -C≡C-, -CH₂-C≡C-, -CH₂-C
 ≡C-CH₂-CH₂-, etc.);

(4) -(CH₂)_{w1}O(CH₂)_{w2}-, -(CH₂)_{w1}S(CH₂)_{w2}-,
 -(CH₂)_{w1}CO(CH₂)_{w2}-, -(CH₂)_{w1}SO(CH₂)_{w2}-,
 25 -(CH₂)_{w1}SO₂(CH₂)_{w2}-, -(CH₂)_{w1}NR^a(CH₂)_{w2}-;

(5) -(CH₂)_{w3}CONR^a(CH₂)_{w4}-, -(CH₂)_{w3}NR^aCO(CH₂)_{w4}-,
 -(CH₂)_{w3}SO₂NR^a(CH₂)_{w4}-, -(CH₂)_{w3}NR^aSO₂(CH₂)_{w4}-,
 -(CH₂)_{w3}COO(CH₂)_{w4}-;

(6) -(CH₂)_{w5}NR^aCONR^a(CH₂)_{w6}-;

30 (7) -(CH₂)_{w7}CONR^a-(CH₂)_{w8}-CONR^{ab}-(CH₂)_{w9}-;
 -CH=CH-CONR^a-; -CH=CH-SO₂NR^a-;

wherein R^a has the same meaning as defined above; R^{ab} has
 the same meaning as R^a; w1 and w2 is an integer of 0 to
 5, and w1 + w2 is 0 to 5; w3 and w4 is an integer of 0 to
 4, and w3 + w4 is 0 to 4; w5 and w6 is an integer of 0 to
 35 3, and w5 + w6 is 0 to 3; w7, w8 and w9 is an integer of

0 to 2, and $w_7 + w_8 + w_9$ is 0 to 2.

The "spacer having a main chain of 1 to 6 atoms" for X, is preferably $-(CH_2)_{w_1}O(CH_2)_{w_2}-$ (symbols have the same meaning as defined above), $-CONR^{8c}-$, $-NR^{8c}CO-$, $-CH=CH-$
 5 $CONR^{8c}-$, $-SO_2NR^{8c}-$ (R^8 is hydrogen atom or C_{1-6} alkyl); more preferably $-CONR^{8c}-$, $-NR^{8c}CO-$, $-CH=CH-CONR^{8c}-$, $-SO_2NR^{8c}-$ (R^8 has the same meaning as defined above); especially preferably $-CONH-$, $-NHCO-$, etc.

10 The "spacer having a main chain of 1 to 6 atoms" for Y, is preferably optionally halogenated bivalent C_{1-6} non-cyclic hydrocarbon groups, $-(CH_2)_{w_3}CONH(CH_2)_{w_4}-$, $-(CH_2)_{w_3}COO(CH_2)_{w_4}-$ (symbols have the same meaning as defined above); more preferably C_{1-3} alkylene (e.g. $-CH_2-$, $-(CH_2)_2-$,
 15 $-(CH_2)_3-$, etc.), $-(CH_2)_{w_3}CONH(CH_2)_{w_4}-$, $-(CH_2)_{w_3}COO(CH_2)_{w_4}-$ (symbols have the same meaning as defined above); especially preferably C_{1-3} alkylene (e.g. $-CH_2-$, $-(CH_2)_2-$, $-(CH_2)_3-$, etc.), etc.

20 As "substituents" and "monocyclic aromatic rings" in "monocyclic aromatic rings which may be condensed with 4 to 8 membered non-aromatic rings, and may have further substituents" for Ar, those exemplified as "substituents" and "cyclic group" in the "cyclic group which may have
 25 substituents" for the above Ar¹ can be used. The number of substituents is, for instance, 1 to 5, preferably 1 to 3. When the number of substituents is 2 or more, each substituents can be the same or different.

The substituents are preferably formyl, optionally
 30 halogenated C_{1-6} alkyl-carbonyl, optionally halogenated C_{1-6} alkylsulfonyl, etc.

Here, as "optionally halogenated C_{1-6} alkyl-carbonyl" and "optionally halogenated C_{1-6} alkylsulfonyl", those exemplified as "substituents" in " C_{7-19} aralkyl which may
 35 have substituents" can be used respectively.

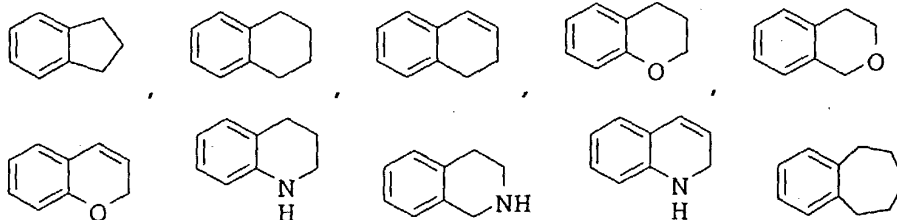
Examples of "4 to 8 membered non-aromatic rings" in the "monocyclic aromatic rings which may be condensed with 4 to 8 membered non-aromatic rings, and may have further substituents" include C_{4-8} monocyclic non-aromatic hydrocarbon rings, 4 to 8 membered monocyclic non-aromatic hetero rings.

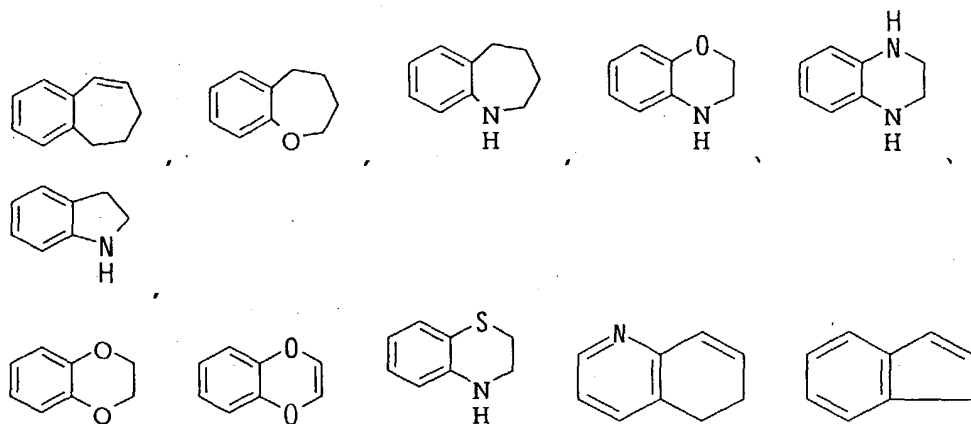
Examples of the " C_{4-8} monocyclic non-aromatic hydrocarbon rings" include C_{4-8} cycloalkane and C_{4-8} cycloalkene. Concrete examples include cyclobutane, cyclopentane, cyclohexane, cycloheptane, cyclooctane, cyclopentene, cyclohexene, cycloheptene. Especially, cyclopentane, cyclohexane, cyclobutane, etc. are preferable.

Examples of the "4 to 8 membered monocyclic non-aromatic hetero rings" include azetidine, pyrrolidine, pyrroline, pyrazolidine, 2- or 3-pyrazoline, imidazoline, piperidine, piperazine, azepine, azokane, oxane, oxine, oxepane, oxazolidine, 2-oxazoline, thiazolidine, 2-thioazoline, morpholine, thiomorpholine.

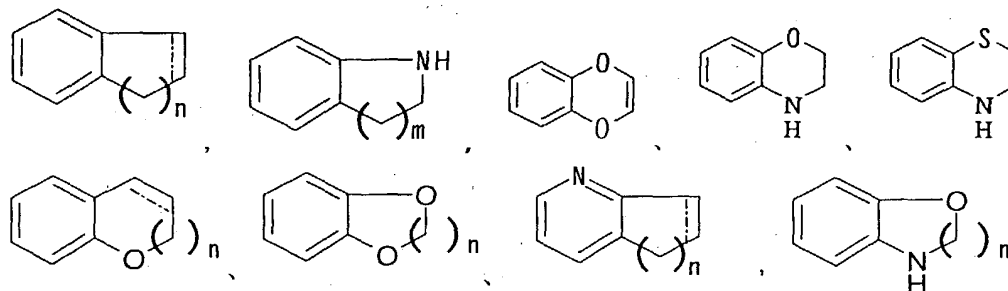
The above "4 to 8 membered non-aromatic rings" may have 1 to 3 substituents at a substitutable position. Examples of such substituents include optionally halogenated C_{1-6} alkyl (e.g., methyl, ethyl, propyl, isopropyl, butyl, isobutyl, sec-butyl, tert-butyl, pentyl, hexyl, etc.), cyano, hydroxy.

Regarding Ar, concrete examples of "monocyclic aromatic rings which may be condensed with 4 to 8 membered non-aromatic rings, and may have further substituents" include



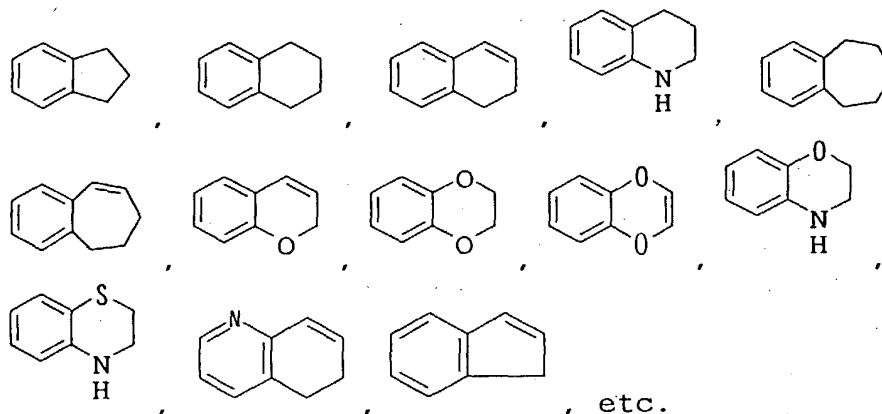


Ar is preferably benzene, pyridine, or rings of the
5 formulae :



wherein ----- is a single bond or double bond; each of m
and n is an integer of 1 to 4.

10 Ar is more preferably benzene, pyridine, rings of the
formulae :



15 As the "hydrocarbon groups which may have
substituents" for R¹ and R², those exemplified as the above R³

can be used.

The "hydrocarbon groups which may have substituents" are preferably "C₁₋₆ alkyl which may have substituents".

Here, examples of "C₁₋₆ alkyl" in the "C₁₋₆ alkyl which
5 may have substituents" include methyl, ethyl, propyl, isopropyl, butyl, isobutyl, sec-butyl, tert-butyl, pentyl, hexyl. Especially, methyl, ethyl, propyl, etc. are preferable.

Examples of "substituents" in the "C₁₋₆ alkyl which may
10 have substituents" include halogen atom (e.g. fluorine, chlorine, bromine, iodine, etc.), C₁₋₃ alkylenedioxy (e.g. methylenedioxy, ethylenedioxy etc.), nitro, cyano, optionally halogenated C₃₋₆ cycloalkyl, optionally halogenated C₁₋₆ alkoxy, optionally halogenated C₁₋₆
15 alkylthio, hydroxy, amino, mono-C₁₋₆ alkylamino (e.g. methylamino, ethylamino, propylamino, isopropylamino, butylamino, etc.), di-C₁₋₆ alkylamino (e.g. dimethylamino, diethylamino, dipropylamino, dibutylamino, ethylmethylamino, etc.), formyl, carboxy, carbamoyl,
20 thiocarbamoyl, optionally halogenated C₁₋₆ alkyl-carbonyl, optionally halogenated C₁₋₆ alkoxy-carbonyl (e.g. methoxycarbonyl, ethoxycarbonyl, propoxycarbonyl, tert-butoxycarbonyl, etc.), mono-C₁₋₆ alkyl-carbamoyl (e.g. methylcarbamoyl, ethylcarbamoyl, etc.), di-C₁₋₆ alkyl-
25 carbamoyl (e.g. dimethylcarbamoyl, diethylcarbamoyl, ethylmethylcarbamoyl etc.), optionally halogenated C₁₋₆ alkylsulfonyl, formylamino, optionally halogenated C₁₋₆ alkyl-carboxamide, C₁₋₆ alkoxy-carboxamide (e.g. methoxycarboxamide, ethoxycarboxamide,
30 propoxycarboxamide, butoxycarboxamide, etc.), C₁₋₆ alkylsulfonylamino (e.g. methylsulfonylamino, ethylsulfonylamino, etc.), C₁₋₆ alkyl-carbonyloxy (e.g. acetoxyl, propanoyloxy, etc.), C₁₋₆ alkoxy-carbonyloxy (e.g. methoxycarbonyloxy, ethoxycarbonyloxy,
35 propoxycarbonyloxy, butoxycarbonyloxy, etc.), mono-C₁₋₆ alkyl-carbamoyloxy (e.g. methylcarbamoyloxy,

ethylcarbamoyloxy, etc.), di-C₁₋₆ alkyl-carbamoyloxy (e.g. dimethylcarbamoyloxy, diethylcarbamoyloxy, etc.), and aromatic groups which may have substituents. The number of substituents is, for instance, 1 to 5, preferably 1 to 3. When the number of substituents is 2 or more, each substituents can be the same or different.

Here, as "optionally halogenated C₃₋₆ cycloalkyl," "optionally halogenated C₁₋₆ alkoxy" and "optionally halogenated C₁₋₆ alkylthio", those exemplified as "substituents" in the above "cyclic group which may have substituents" can be used.

As "optionally halogenated C₁₋₆ alkyl-carbonyl," "optionally halogenated C₁₋₆ alkylsulfonyl" and "optionally halogenated C₁₋₆ alkyl-carboxamide", those exemplified as "substituents" in the above "C₇₋₁₉ aralkyl which may have substituents" can be used.

As "substituents" and "aromatic groups" in the "aromatic groups which may have substituents", those exemplified as "substituents" and "aromatic groups" in the "cyclic group which may have substituents" for the above Ar¹ can be used. The number of substituents is, for instance, 1 to 5, preferably 1 to 3. When the number of substituents is 2 or more, each substituents can be the same or different.

Examples of "nitrogen-containing hetero rings" in the "nitrogen-containing hetero rings which may have substituents" formed by R¹ and R² together with the adjacent nitrogen atom, include 3 to 8 membered nitrogen-containing hetero rings which contain at least one nitrogen atom in addition to carbon atoms, and which may further contain 1 to 3 hetero atoms selected from nitrogen, sulfur and oxygen atom. Concrete examples include aziridine, azetidine, morpholine, thiomorpholine, piperidine, piperazine, pyrrolidine, hexamethyleneimine, heptamethyleneimine, hexahydropyrimidine, 1,4-diazepan, 4,5-dihydroimidazole, and their unsaturated cyclic amines (e.g.

1,2,5,6-tetrahydropyridine, etc.) can be mentioned. Especially, morpholine, piperidine, piperazine, pyrrolidine.

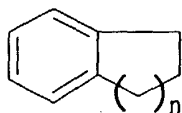
As "substituents" in the "nitrogen-containing hetero rings which may have substituents", for instance, those exemplified as "substituents" in the above "5 to 7 membered saturated cyclic amino which may have substituents" can be used. The number of substituents is, for instance, 1 to 5, preferably 1 to 3. When the number of substituents is 2 or more, each substituents can be the same or different.

R^1 and R^2 are preferably C_{1-6} alkyl, more preferably methyl, ethyl, propyl, etc.

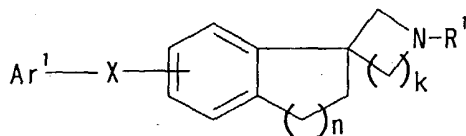
Also, it is preferable that R^1 and R^2 , together with the adjacent nitrogen atom, form piperidino, pyrrolidin-1-yl, piperazin-1-yl etc.

And, it is preferable that at least one of R^1 and R^2 is C_{1-6} alkyl which may have substituents. It is especially preferable that both R^1 and R^2 is C_{1-6} alkyls which may have substituents.

R^2 can form a spiro ring together with Ar. For instance, Ar is a ring of the formula :



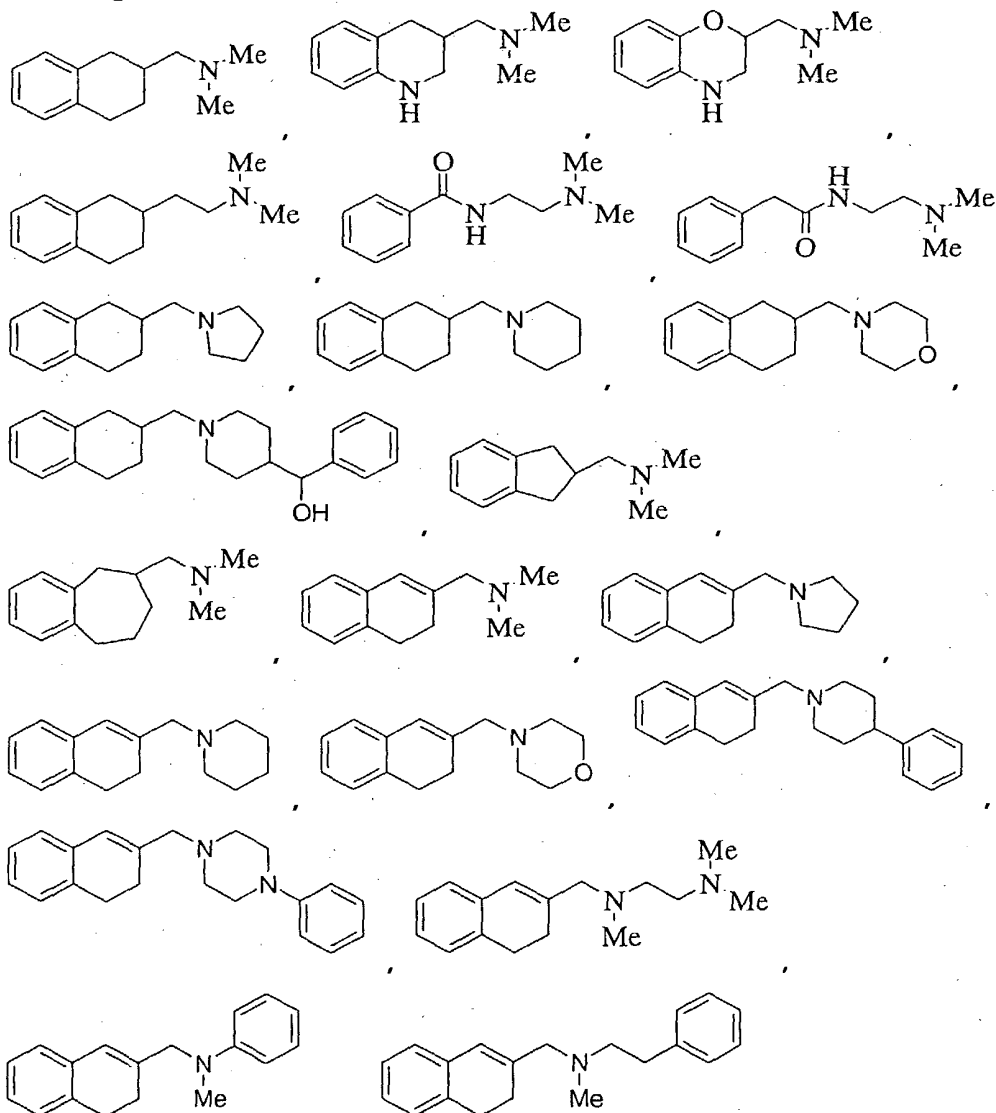
wherein n is an integer of 1 to 4; and Y is methylene; R^2 can form a spiro ring together with Ar. Examples of the spiro ring include

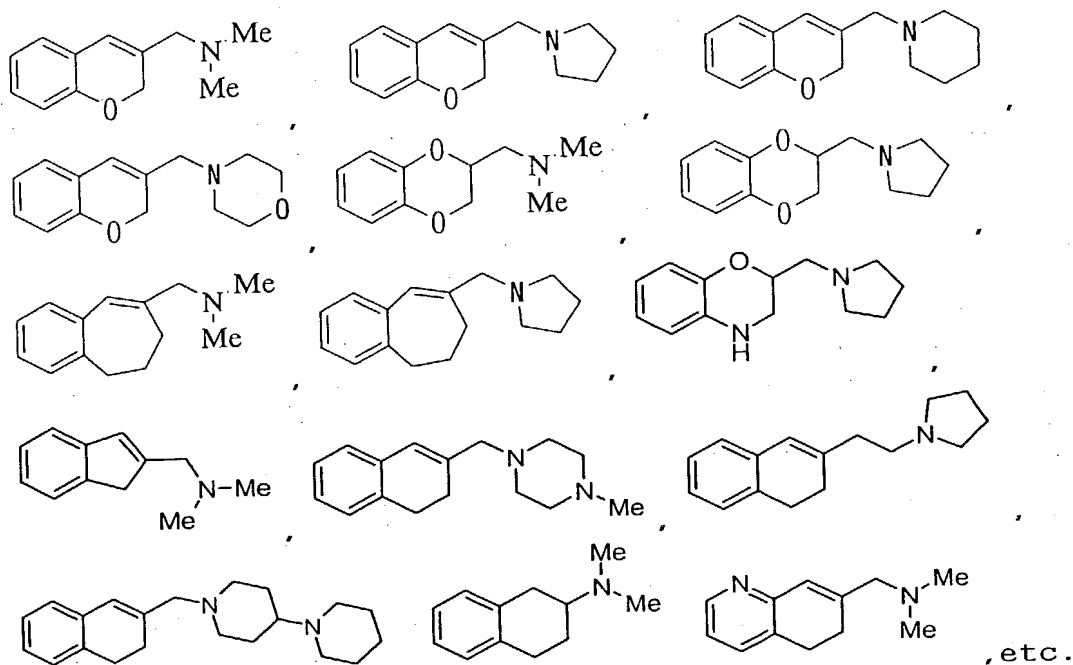


wherein k (ring Ar and N are connected by $-(CH_2)_k-$) is an integer of 1 to 4; and other symbols have the same meaning as defined above.

10

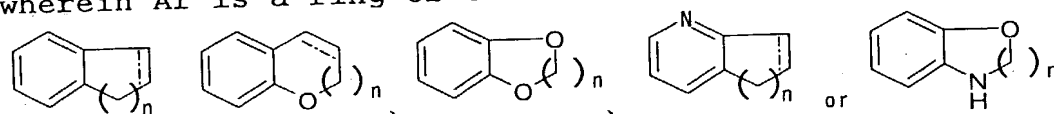
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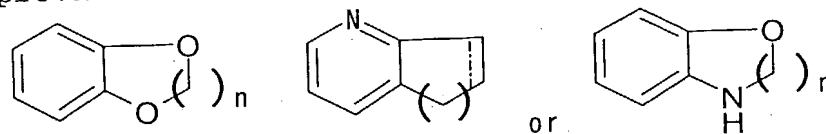


5

Among the compounds of the formula (I), a compound wherein Ar is a ring of the formula :



- wherein ----- is a single bond or double bond, n is an integer of 1 to 4, and each ring may have substituents;
 10 X is $-\text{CONR}^{\text{sc}}-$, $-\text{NR}^{\text{sc}}\text{CO}-$, $-\text{CH}=\text{CH}-\text{CONR}^{\text{sc}}-$ or $-\text{SO}_2\text{NR}^{\text{sc}}-$ where R^{sc} is hydrogen atom or C_{1-6} alkyl;
 Y is a spacer having a main chain of 1 to 6 atoms;
 provided that Ar is a ring of the formulae :



15

- wherein symbols have the same meanings as defined above, and each ring may have substituents, when X is $-\text{SO}_2\text{NH}-$; and provided that Ar^1 is not biphenyl which may be substituted; when X is $-\text{CONH}-$ and Ar is any one of
 20 benzopyran, dihydrobenzopyran, dihydrobenzoxazine, dihydrobenzoxazole or tetrahydrobenzoxazepine;

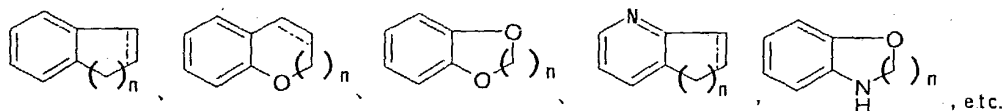
(excluding N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]-4-biphenylcarboxamide);

namely compound of the formula (I') (excluding N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]-4-

5 biphenylcarboxamide) is a novel compound.

Preferred examples of compound of the formula (I') include compound of the formula (I'-1), (I'-2), (I'-3), (I'-4), (I'-5), (I'-6), (I'-7), (I'-8), (I'-9) or (I'-10).

10 In the above formulae (I'), (I'-1), (I'-2), (I'-3), (I'-4), (I'-5), (I'-6), (I'-7), (I'-8), (I'-9) and (I'-10), a ring of the formula :



wherein symbols have the same meanings as above, may have further 1 to 3 substituents at substitutable positions.

15 Examples of such substituents include "substituents" exemplified in the above Ar. Especially, preferred are formyl, optionally halogenated C_{1-6} alkyl-carbonyl, optionally halogenated C_{1-6} alkylsulfonyl, optionally
20 halogenated C_{1-6} alkyl (e.g., methyl, ethyl, propyl, isopropyl, butyl, isobutyl, sec-butyl, tert-butyl, pentyl, hexyl, etc.), cyano, hydroxy, etc.

25 Examples of salts of compound (I) or (I') include salts with inorganic bases, ammonium salts, salts with organic bases, salts with inorganic acids, salts with organic acids, and salts with basic or acidic amino acids.

30 Preferred examples of salts with inorganic bases include alkali metal salts such as sodium salts and potassium salts; alkaline earth metal salts such as calcium salts, magnesium salts, barium salts; and aluminum salts.

Preferred examples of salts with organic bases include salts with trimethylamine, triethylamine, pyridine, picoline, ethanolamine, diethanolamine, triethanolamine,

dicyclohexylamine, N,N-dibenzylethylenediamine.

Preferred examples of salts with inorganic acids include salts with hydrochloric acid, hydrobromic acid, nitric acid, sulfuric acid, phosphoric acid.

5 Preferred examples of salts with organic acids include salts with formic acid, acetic acid, trifluoroacetic acid, fumaric acid, oxalic acid, tartaric acid, maleic acid, citric acid, succinic acid, malic acid, methanesulfonic acid, benzenesulfonic acid, and p-toluenesulfonic acid,
10 3-chlorobenzoic acid.

Preferred examples of salts with basic amino acids include salts with arginine, lysine, ornithine. Preferred examples of salts with acidic amino acids include salts with aspartic acid, glutamic.

15 Among these salts, pharmaceutically acceptable salts are preferable. For instance, when compound (I) or (I') possesses an acidic functional group, it can form an inorganic salt such as an alkali metal salt (e.g., sodium salt, potassium salt, etc.), an alkaline earth metal salt
20 (e.g. calcium salt, magnesium salt, barium salt, etc.), and an ammonium salt. When compound (I) or (I') possesses a basic functional group, it can form an inorganic salt such as hydrochloride, sulfate, phosphate, hydrobromate, etc.; or an organic salt such as acetate, maleate, fumarate,
25 succinate, methanesulfonate, p-toluenesulfonate, citrate and tartrate.

Compounds (I) and (I') (hereinafter also abbreviated as a compound of the invention) can be either anhydrides or hydrates. A hydrate may have 0.5 to 3 water molecules.

30 In addition, a compound of the invention can be labeled using isotopes (e.g. ^3H , ^{14}C , and ^{35}S , etc.).

When a compound of the invention contains optical isomers, stereoisomers, regio isomers, rotational isomers,
35 these are included as a compound of the invention, and each of them can be obtained as a single substance by per se known

5 The optical isomers can be produced using per se known methods. Concretely, the optical isomer can be obtained by using an optically active synthetic intermediate, or subjecting the racemic mixture of the final product to optical resolution in accordance with common method.

1) Fractional recrystallization method

2) Chiral column method

This method comprises subjecting a racemate or its salt to a column for separating an optical isomer (chiral column) for separation. For instance, in the case of liquid chromatography, an optical isomer mixture is added to the chiral column such as ENANTIO-OVM [produced by Tosol] or CHIRAL series [produced by Daicel], which is developed using water, various buffer solutions (e.g. phosphate buffer), organic solvents (e.g. ethanol, methanol, isopropanol, acetonitrile, trifluoroacetic acid, diethylamine, etc.) as single or mixed solutions, and the optical isomers are separated. Also, in the case of gas chromatography, for instance, separation is conducted using a chiral column such as CP-Chirasil-DeX (produced by

G.L.Science Co.).

3) Diastereomer method

In this method, a racemic mixture is subjected to a chemical reaction with an optically active reagent to give
5 a diastereomer mixture, which is separated into a single substance by an ordinary separation means (e.g. fractional recrystallization, chromatography method, etc.). This single substance is subjected to removal of the optically active reagent part using chemical processing such as a
10 hydrolysis reaction. For instance, when a compound of the invention possesses hydroxy or primary or secondary amino in its molecule, this compound is subjected to a condensation reaction with an optically active organic acid (e.g. MTPA [α -methoxy- α -(trifluoromethyl)phenylacetic
15 acid], (-)-menthoxyacetic acid, etc.), to give the diastereomer in an ester form or an amide form, respectively. On the other hand, when a compound of the invention possesses carboxylic acid group, this compound is subjected to a condensation reaction with an optically
20 active amine or alcohol reagent, to give the diastereomer in an amide form or an ester form, respectively. The separated diastereomer can be converted to an optical isomer of the original compound, by applying acidic hydrolysis or basic hydrolysis.

25 A prodrug of compound (I') is a compound which is converted to compound (I') by reactions involving enzymes and gastric acid, etc. under physiological conditions in the living body; in other words, a compound that is changed into compound (I') by enzymatically-caused oxidation,
30 reduction and hydrolysis, and a compound that is changed into compound (I') by hydrolysis caused by gastric acid.

Examples of the prodrugs of compound (I') include compounds in which amino groups of compound (I') have been acylated, alkylated, or phosphorylated [e.g. compounds in
35 which amino groups of compound (I') have been eicosanoylated, aranylated, pentylaminocarbonylated,

(5-methyl-2-oxo-1,3-dioxolen-4-yl)methoxycarbonylated, tetrahydrofuranylated, pyrrolidylmethylated, pivaloyloxymethylated, tert-butylated, etc.]; compounds in which hydroxyl groups of compound (I') have been acylated, alkylated, phosphorylated, borated (e.g. compounds in which hydroxyl groups of compound (I') have been acetylated, palmitoylated, propanoylated, pivaloylated, succinylated, fumarilated, alanilated, dimethylaminomethylcarbonylated, etc.); compounds in which carboxyl groups of compound (I') have been esterified or amidated [e.g. compounds in which carboxyl groups of compound (I') have been ethylesterified, phenylesterified, carboxylmethylesterified, dimethylaminomethylesterified, pivaloyloxymethylesterified, ethoxycarbonyloxyethylesterified, phthalidylesterified, (5-methyl-2-oxo-1,3-dioxolen-4-yl)methylesterified, cyclohexyloxycarbonylethylesterified, or methylamidated, etc.]. These compounds can be produced from compound (I') using per se known methods.

Also, a prodrug of compound (I') can be a compound which is changed to compound (I') by physiological conditions, as described in pages 163 to 198 of Molecular Design, Volume 7, "Development of Drugs," published in 1990 by Hirokawa Shoten.

A compound of the invention can be produced in accordance with per se known methods such as methods described in WO9838156, WO9532967, and EP-A533266, etc., or analogous methods thereto.

For instance, a compound of the invention can be produced in accordance with [Production method 1] to [Production method 6] which are described in detail below, or analogous methods thereto.

Compounds (II) to (XI) used as raw materials, can be used in the form of salts. As such salts, those exemplified

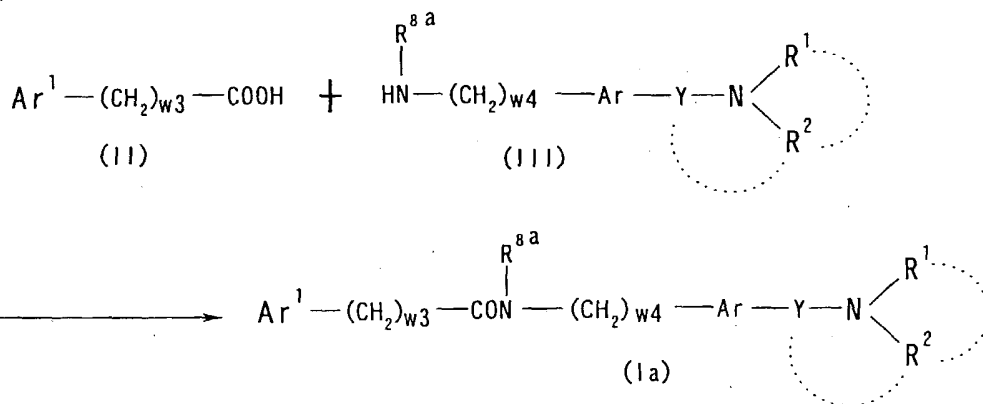
as salts of the above compound (I) or (I') can be used.

In the following [Production method 1] to [Production method 6], when an alkylation reaction, a hydrolysis reaction, an amination reaction, an esterification reaction, an amidation reaction, an esterification reaction, an etherification reaction, an oxidation reaction, a reduction reaction, etc. are carried out, these reactions are carried out in accordance with per se known methods. Examples of such methods include the methods described in Organic Functional Group Preparations, Second Edition, Academic Press, Inc., published in 1989; Comprehensive Organic Transformations, VCH Publishers Inc., published in 1989, etc.

[Production method 1]

Compound (Ia) having $-(CH_2)_{w3}CONR^{8a}(CH_2)_{w4}-$ for X in formula (I), is produced, for instance, by the following amidation reaction.

(Amidation reaction)



wherein R^{8a} is hydrogen atom or an optionally halogenated C_{1-6} alkyl; other symbols have the same meanings as defined above.

As the "optionally halogenated C_{1-6} alkyl", those exemplified as "substituents" in the above "cyclic group which may have substituents" can be used.

The "amidation reaction" includes the following

"method using a dehydration and condensation agent" and
"method using a reactive derivative of carboxylic acid".

i) Method using a dehydration and condensation agent

5 Compound (III), 1 to 5 equivalents of compound (II),
and 1 to 2 equivalents of a dehydration and condensation
agent are reacted in an inert solvent. If necessary, the
reaction can be carried out with the coexistence of 1 to
10 1.5 equivalents of 1-hydroxybenzotriazole (HOBT) and (or)
catalytic quantity to 5 equivalents of a base.

Examples of the "dehydrating and condensation agent"
include dicyclohexylcarbodiimide (DCC), 1-ethyl-3-(3-
dimethylaminopropyl)carbodiimide hydrochloride (WSC).
WSC is particularly preferable.

15 Examples of the "inert solvent" include nitrile
solvents (preferably acetonitrile), amide solvents
(preferably DMF), halogenated hydrocarbon solvents
(preferably dichloromethane), ether solvents (preferably
THF). Two or more kinds of these can be mixed in an
20 appropriate ratio for use.

Examples of the "base" include

1) for instance, strong bases such as hydrides of
alkali metals or alkaline earth metals (e.g. lithium
25 hydride, sodium hydride, potassium hydride, calcium
hydride, etc.), amides of alkali metals or alkaline earth
metals (e.g. lithium amide, sodium amide, lithium
diisopropylamide, lithium dicyclohexylamide, lithium
hexamethyldisilazide, sodium hexamethyldisilazide,
30 potassium hexamethyldisilazide, etc.), lower alkoxides of
alkali metals or alkaline earth metals (e.g. sodium
methoxide, sodium ethoxide, potassium tert-butoxide,
etc.);

2) for instance, inorganic bases such as hydroxides
35 of alkali metals or alkaline earth metals (e.g. sodium
hydroxide, potassium hydroxide, lithium hydroxide, barium

hydroxide, etc.), carbonates of alkali metals or alkaline earth metals (e.g. sodium carbonate, potassium carbonate, cesium carbonate, etc.) and hydrogencarbonates of alkali metals or alkaline earth metals (e.g. sodium
5 hydrogencarbonate, potassium hydrogencarbonate, etc.); and

3) for instance, amines such as triethylamine, diisopropylethylamine, N-methylmorpholine, dimethylaminopyridine, DBU (1,8-
10 diazabicyclo[5.4.0]undec-7-en), DBN (1,5-diazabicyclo[4.3.0]non-5-en); for instance, organic bases such as basic heterocyclic compounds of pyridine, imidazole, 2,6-lutidine, etc.

Among the above bases, triethylamine, 4-
15 dimethylaminopyridine, etc., are preferable.

Reaction temperature is usually room temperature (0°C to 30°C, hereafter the same). Reaction time is, for instance, 10 to 24 hours.

ii) Method using a reactive derivative of carboxylic acid.
20 A reactive derivative of compound (II) and 1 to 5 equivalents (preferably 1 to 3 equivalents) of compound (III) are reacted in an inert solvent. If necessary, the reaction can be carried out with the coexistence of 1 to 10 equivalents, preferably 1 to 3 equivalents of a base.

25 Examples of the "reactive derivative" of compound (II) include acid halides (e.g., acid chloride, acid bromide, etc.), mixed acid anhydrides (e.g. acid anhydrides with C₁₋₆ alkyl-carboxylic acid, C₆₋₁₀ aryl-carboxylic acid or C₁₋₆ alkylcarbonate), active esters (e.g. esters with phenol
30 which may have substituents, 1-hydroxybenzotriazole or N-hydroxysuccinimide, etc.).

Examples of the "substituents" in the "phenol which may have substituents" include halogen atom (e.g. fluorine, chlorine, bromine, iodine, etc.), nitro, optionally
35 halogenated C₁₋₆ alkyl, optionally halogenated C₁₋₆ alkoxy. The number of substituents is, for instance, 1 to 5.

As the "optionally halogenated C₁₋₆ alkyl" and "optionally halogenated C₁₋₆ alkoxy", those exemplified as "substituents" in the above "cyclic group which may have substituents" can be used.

5 Concrete examples of "phenol which may have substituents" include phenol, pentachlorophenol, pentafluorophenol, p-nitrophenol. The reactive derivative is, preferably, an acid halide.

10 Examples of the "inert solvent" include ether solvents, halogenated hydrocarbon solvents, aromatic solvents, nitrile solvents, amide solvents, ketone solvents, sulfoxide solvents, and water. Two or more kinds of these can be mixed in an appropriate ratio for use.

Especially, acetonitrile, THF, dichloromethane, 15 chloroform, etc. are preferable.

As the "base", the same as above are used. The base is preferably sodium hydride, potassium carbonate, sodium carbonate, sodium hydroxide, potassium hydroxide, sodium hydrogencarbonate, potassium hydrogencarbonate, 20 triethylamine, pyridine, etc.

Reaction temperature is usually -20°C to 50°C, preferably room temperature. Reaction time is usually 5 minutes to 40 hours, preferably 1 to 18 hours.

Compound (III) can be produced by per se known methods. 25 For instance, 6-amino-2-(N,N-dimethylamino)methyltetraline or its salt can be produced in accordance with the methods described in WO9838156.

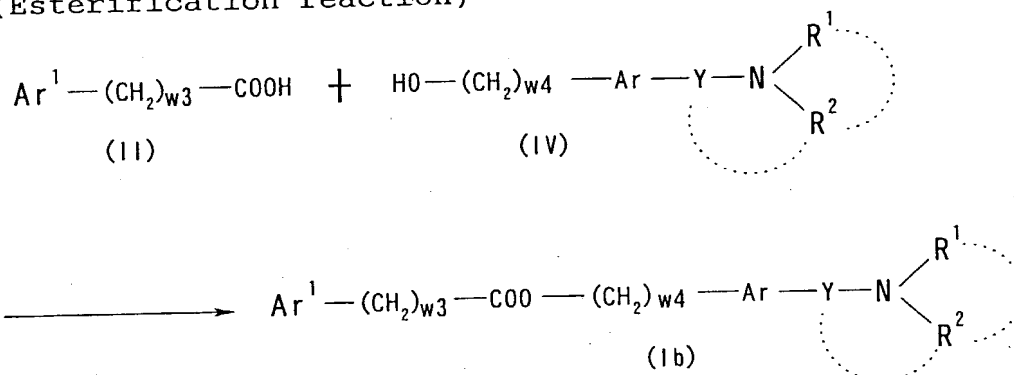
Also, 6-amino-2,3-dihydro-1-(2-dimethylaminoethyl)-1H-indole, 6-amino-3,4-dihydro-4-(2-dimethylaminoethyl)- 30 2H-1,4-benzoxazine, etc., can be produced in accordance with the methods described in WO9532967.

The above "method using a reactive derivative of carboxylic acid" can be also adopted when producing a corresponding sulfonamide derivative or sulfinamide 35 derivative, from the sulfonic acid of the formula :
$$\text{Ar}^1-(\text{CH}_2)_{w_3}-\text{SO}_2\text{OH}$$
 (symbols have the same meanings as defined

above), or the sulfinic acid of the formula : $\text{Ar}^1-(\text{CH}_2)_{w3}-\text{SOOH}$
(symbols have the same meanings as defined above).

[Production method 2]

- 5 Compound (Ib) having $-(\text{CH}_2)_{w3}-\text{COO}(\text{CH}_2)_{w4}-$ for X in the
formula (I), can be produced by the following
esterification reaction.
(Esterification reaction)



- 10 wherein symbols have the same meanings as defined above.

A reactive derivative of compound (II) and 1 to 5
equivalents (preferably 1 to 3 equivalents) of compound
(IV) is reacted in an inert solvent. Usually, this reaction
is carried out with the coexistence of 1 to 10 equivalents,
15 preferably 1 to 3 equivalents of a base.

As the reactive derivative of compound (II), the same
as above is used. Especially, an acid halide is preferable.

20 Examples of the "inert solvent" include ether
solvents, halogenated hydrocarbon solvents, aromatic
solvents, nitrile solvents, amide solvents, ketone
solvents, sulfoxide solvents. Two or more kinds of these
can be mixed in an appropriate ratio for use. Especially,
acetonitrile, dichloromethane, chloroform, etc. are
preferable.

25 As the "base", the same one as above can be used. The
base is preferably sodium hydride, potassium carbonate,
sodium carbonate, sodium hydroxide, potassium hydroxide,
sodium hydrogencarbonate, potassium hydrogencarbonate,

triethylamine, pyridine, etc.

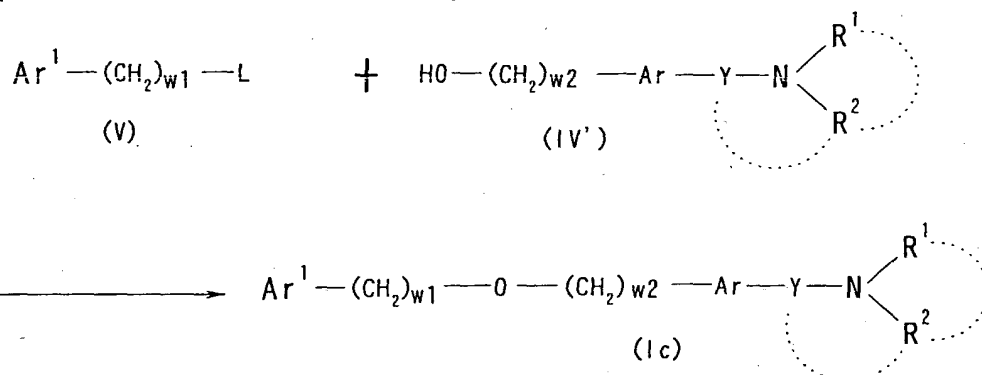
Reaction temperature is usually -20°C to 50°C , preferably room temperature. Reaction time is usually 5 minutes to 40 hours, preferably 1 to 18 hours.

5

[Production method 3]

Compound (Ic) having $-(\text{CH}_2)_{w1}\text{O}(\text{CH}_2)_{w2}-$ for Y in the formula (I), can be produced by, for instance, the following etherification reaction.

10 (Etherification reaction)



wherein L is a leaving group, and other symbols have the same meanings as defined above.

15 Examples of the "leaving group" for L include halogen atom (e.g. chlorine, bromine, iodine, etc.), optionally halogenated C_{1-6} alkylsulfonyloxy (e.g. methanesulfonyloxy, ethanesulfonyloxy, trifluoromethanesulfonyloxy, etc.), C_{6-10} arylsulfonyloxy which may have substituents, hydroxy.

20 Examples of the "substituents" in the " C_{6-10} arylsulfonyloxy which may have substituents" include halogen atom (e.g. chlorine, bromine, iodine, etc.), optionally halogenated C_{1-6} alkyl, C_{1-6} alkoxy. The number of substituents is, for instance, 1 to 3. Concrete examples
25 of the C_{6-10} arylsulfonyloxy which may have substituents" include benzenesulfonyloxy, p-toluenesulfonyloxy, 1-naphthalenesulfonyloxy, 2-naphthalenesulfonyloxy.

The "leaving group" is preferably halogen atom (e.g.

chlorine, bromine, iodine, etc.), methanesulfonyloxy, trifluoromethanesulfonyloxy, p-toluenesulfonyloxy.

Compound (IV') and about 1 to 5 equivalents
5 (preferably 1 to 2 equivalents) of compound (V) are reacted in inert solvent, with the coexistence of base.

As the "base", the same one as above can be used. The base is preferably potassium carbonate, sodium hydrogencarbonate, triethylamine, N-methylmorpholine,
10 pyridine, etc. The amount of the base used is usually about 1 to 5 equivalents relative to compound (V).

Examples of the "inert solvent" include alcohol solvents, ether solvents, halogenated hydrocarbon solvents, aromatic solvents, nitrile solvents, amide
15 solvents, ketone solvents, sulfoxide solvents, water. Two or more kinds of these can be mixed in an appropriate ratio for use. Especially, acetonitrile, N,N-dimethylformamide (DMF), acetone, ethanol, pyridine, etc., are preferable.

Reaction temperature is about -20°C to 100°C,
20 preferably room temperature to 80°C. Reaction time is, for instance, 5 hours to 1 day.

In the above production method, when the leaving group is hydroxy, Mitsunobu reaction can usually be used. In the Mitsunobu reaction, compound (V) and 0.5 to 5 equivalents
25 (preferably 1 to 1.5 equivalents) of compound (IV') are reacted in inert solvent with the coexistence of 0.5 to 5 equivalents (preferably 1 to 1.5 equivalents) of ethyl acetyldicarboxylate.

Examples of the inert solvent include ether solvents,
30 halogenated hydrocarbon solvents, aromatic solvents, nitrile solvents, amide solvents, ketone solvents, sulfoxide solvents. Two or more kinds of these can be mixed in an appropriate ratio for use. Especially, acetonitrile, dichloromethane, chloroform, etc. are
35 preferable.

Reaction temperature is usually -20°C to 50°C,

preferably room temperature. Reaction time is usually 5 minutes to 40 hours, preferably 1 to 18 hours.

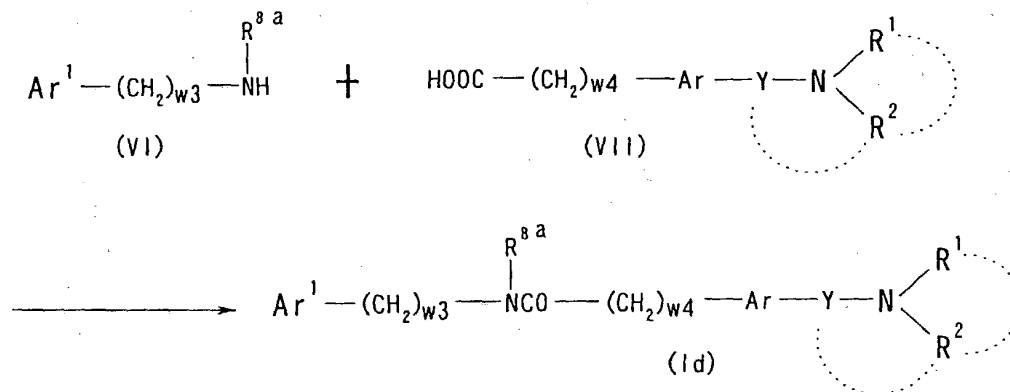
Compound (IV') can be produced by per se known methods.

For instance, 3-(N,N-dimethylamino)methyl-1,2,3,4-tetrahydro-7-quinolinol, 2-(N,N-dimethylamino)methyl-6-hydroxytetralin, 6-hydroxy-2-piperidinomethyltetralin, 2-[2-(N,N-dimethylamino)ethyl]-6-hydroxytetralin, 2-(N,N-dimethylamino)methyl-7-hydroxytetralin, 6-hydroxy-2-(N-methylamino)methyltetralin, etc., can be produced in accordance with the methods described in WO9838156.

[Production method 4]

Compound (Id) having $-(CH_2)_wNR^{8a}CO(CN_2)_w-$ for X in the formula (I), can be produced, for instance, by the following amidation reaction.

(Amidation reaction)



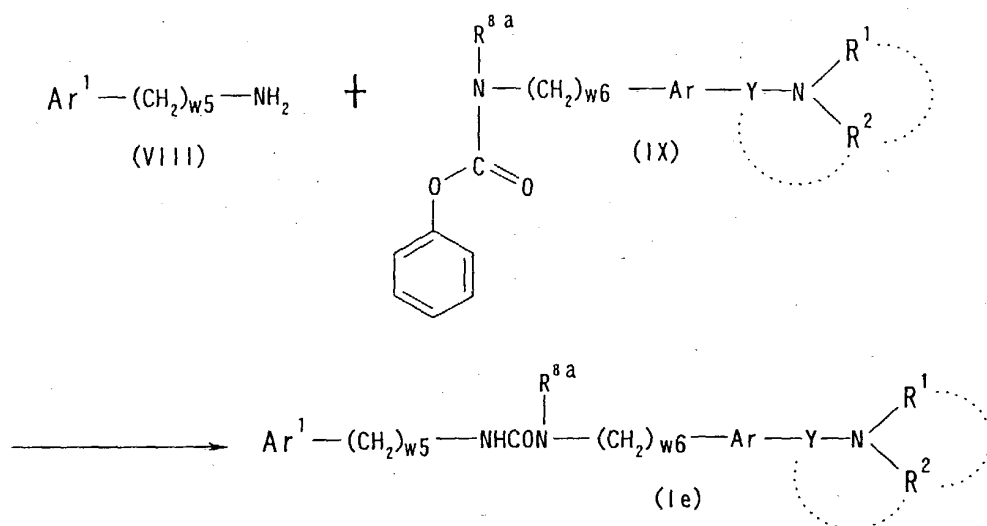
wherein symbols have the same meanings as defined above.

This Production method is carried out in accordance with the above Production method 1.

[Production method 5]

Compound (Ie) having $-(CH_2)_wNHCONR^{8a}(CN_2)_w-$ for X in the formula (I), can be produced, for instance, by the following urea reaction.

(Urea reaction)



wherein symbols have the same meanings as defined above.

Compound (IX) and 1 to 5 equivalents (preferably 1 to 1.5 equivalents) of compound (VIII) is reacted in an inert solvent with the coexistence of a base.

As the "base", the same one as above can be used. The base is preferably potassium carbonate, sodium carbonate, sodium hydroxide, potassium hydroxide, sodium hydrogencarbonate, potassium hydrogencarbonate, triethylamine, pyridine, etc.

Examples of the "inert solvent" include alcohol solvents, ether solvents, halogenated hydrocarbon solvents, aromatic solvents, nitrile solvents, amide solvents, ketone solvents, sulfoxide solvents, water. Two or more kinds of these can be mixed in an appropriate ratio for use. Especially, acetonitrile, DMF, acetone, ethanol, pyridine, etc. are preferable.

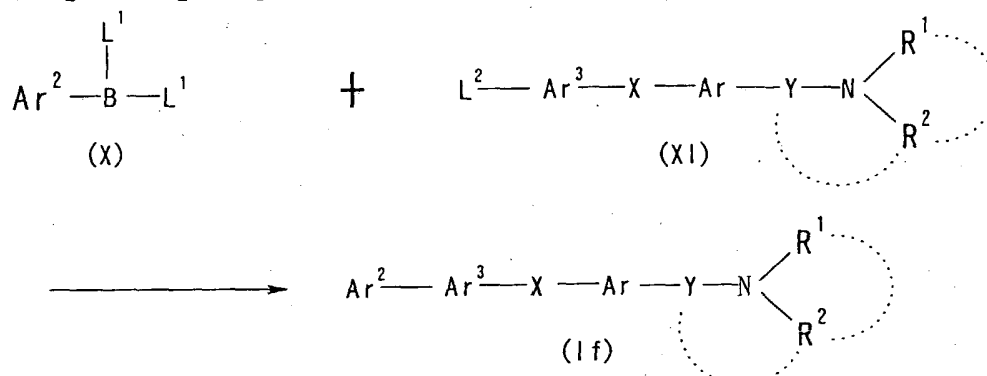
Reaction temperature is usually -20°C to 100°C , preferably room temperature to 80°C . Reaction time is, for instance, 0.5 hour to 1 day.

[Production method 6]

Compound (If) having, for Ar^1 , a ring assembly aromatic

group ($\text{Ar}^2\text{-Ar}^3$) which may have substituents in the formula (I), can be produced by, for instance, the following aryl-coupling reaction.

(Aryl-coupling reaction)



5

wherein Ar^2 and Ar^3 are monocyclic aromatic groups or condensed aromatic groups, each of which may have substituents; L^1 is hydroxy or C_{1-6} alkyl; L^2 is halogen (preferably chlorine, bromine) or trifluoromethanesulfonyloxy; other symbols have the same meanings as defined above.

10

As "substituents", "monocyclic aromatic groups" and "condensed aromatic groups" in the "monocyclic aromatic groups or condensed aromatic groups, each of which may have substituents" for Ar^2 and Ar^3 , those exemplified as the above Ar^1 can be used. Especially, it is preferable that both of Ar^2 and Ar^3 are phenyl groups which may have substituents, and $\text{Ar}^2\text{-Ar}^3$ is biphenyl which may have substituents.

15

The aryl-coupling reaction can be carried out in accordance with per se known methods such as the method described in Acta. Chemica Scandinavia, pp. 221-230, 1993, or methods analogous thereto.

20

Compound (X) and 1 to 3 equivalents (preferably 1 to 1.5 equivalents) of compound (XI) are reacted in an inert solvent in the presence of a base and a transition metal catalyst.

25

As the base, the same one as above can be used. The

base is preferably sodium carbonate, sodium hydrogencarbonate, etc.

The amount of the "base" used is, for instance, about 1 to 10 equivalents relative to compound (XI).

5 Examples of the "transition metal catalyst" include palladium catalyst, nickel catalyst. Examples of the "palladium catalyst" include tetrakis(triphenylphosphine)palladium (O), palladium acetate, bis (triphenylphosphine) palladium (II) chloride, 10 palladium-carbon. Examples of the "nickel catalyst" include tetrakis(triphenylphosphine) nickel (O).

The amount of the "transition metal catalyst" used is about 0.01 to .1 equivalent, preferably about 0.01 to 0.5 equivalent, relative to compound (XI).

15 Reaction temperature is room temperature to 150°C, preferably about 80°C to 150°C. Reaction time is, for instance, about 1 to 48 hours.

Examples of the "inert solvent" include water, alcohol solvents, aromatic solvents. Two or more kinds of these 20 can be mixed in an appropriate ratio for use. Especially, a single solvent such as water, ethanol and toluene; or a mixed solvent of two or more kinds of these is preferable.

Examples of the above "alcohol solvents" include methanol, ethanol, isopropanol, tert-butanol.

25 Examples of the above "ether solvents" include diethylether, tetrahydrofuran (THF), 1,4-dioxane, 1,2-dimethoxyethane.

Examples of the above "halogenated hydrocarbon solvents" include dichloromethane, chloroform, 1,2- 30 dichloroethane, carbon tetrachloride.

Examples of the above "aromatic solvents" include benzene, toluene, xylene, pyridine.

Examples of the above "hydrocarbon solvents" include hexane, pentane, cyclohexane.

35 Examples of the above "amide solvents" include N,N-dimethylformamide (DMF), N,N-dimethylacetamide, N-

Examples of the above "ketone solventd" include acetone, methylethylketone.

Examples of the above "nitrile solvents" include acetonitrile, propionitrile.

In a compound of the invention thus obtained, the intramolecular functional group can be converted to a desired functional group by combining per se known chemical reactions. Examples of the chemical reactions include oxidation reaction, reduction reaction, alkylation reaction, hydrolysis reaction, amination reaction, esterification reaction, aryl-coupling reaction, deprotection reaction.

In each of the above reactions, when the raw material compounds possess amino, carboxy, hydroxy, and/or carbonyl as substituents, protecting groups which are generally used in peptide chemicals, etc., can be introduced into these groups, and the desired compound can be obtained by removing the protecting groups after the reaction if necessary.

Examples of the protecting group for amino include formyl, C₁₋₆ alkyl-carbonyl (e.g. acetyl, propionyl, etc.), C₁₋₆ alkoxy-carbonyl (e.g. methoxycarbonyl, ethoxycarbonyl, tert-butoxycarbonyl, etc.), benzoyl, C₇₋₁₀ aralkyl-carbonyl (e.g. benzylcarbonyl, etc.), C₇₋₁₄ aralkyloxy-carbonyl (e.g. benzyloxycarbonyl, 9-fluorenylmethoxycarbonyl, etc.), trityl, phthaloyl, N,N-dimethylaminomethylene, silyl (e.g. trimethylsilyl, triethylsilyl, dimethylphenylsilyl, tert-butyl dimethylsilyl, tert-butyl diethylsilyl, etc.), C₂₋₆ alkenyl (e.g. 1-allyl, etc.) . These groups may be substituted by 1 to 3 of halogen atom (e.g. fluorine, chlorine, bromine, iodine, etc.), C₁₋₆ alkoxy (e.g. methoxy, ethoxy, propoxy, etc.) or nitro, etc.

Examples of the protecting group for carboxy include

Removal of the above protecting groups can be carried out in accordance with per se known methods such as those described in Protective Groups in Organic Synthesis, published by John Wiley and Sons (1980). For instance, the methods using acid, base, ultraviolet light, hydrazine, phenylhydrazine, sodium N-methyldithiocarbamate, tetrabutylammonium fluoride, palladium acetate, trialkylsilyl halide (e.g. trimethylsilyl iodide, trimethylsilyl bromide, etc.), and a reduction method, etc. can be used.

5 It is also possible to isolate and purify the raw material compounds of a compound of the invention, or their salts using the same known methods as above, but they can also be used as raw materials in the next process as a reaction mixture without being isolated.

Also, a compound of the invention is low in toxicity, and
15 is excellent in oral absorbency and intracerebral
transitivity.

Here, examples of the diseases caused by MCH include obesity (e.g. malignant mastocytosis, exogenous obesity, hyperinsular obesity, hyperplasmic obesity, hypophyseal adiposity, hypoplasmic obesity, hypothyroid obesity, hypothalamic obesity, symptomatic obesity, infantile obesity, upper body obesity, alimentary obesity, hypogonadal obesity, systemic mastocytosis, simple obesity, central obesity, etc.), hyperphagia, emotional disorders, reproductive function disorders, memory disorders, dementia, hormonal disorders.

A compound of the invention is also useful as an agent
35 for preventing or treating lifestyle diseases such as
diabetes, diabetic complications (e.g. diabetic

retinopathy, diabetic neuropathy, diabetic nephropathy, etc.), arteriosclerosis, and gonitis.

Further, a compound of the invention is useful as an anorectic agent.

5 A MCH antagonist and a pharmaceutical composition of the invention can be used in combination with an alimentary therapy (e.g., alimentary therapy for diabetes) and exercise.

10 A MCH antagonist and a pharmaceutical composition of the invention can be produced by subjecting compound (I) or compound (I') respectively, as it is, or together with a pharmacologically acceptable carrier, to pharmaceutical manufacturing process in accordance with a per se known means.

15 Here, examples of the pharmacologically acceptable carriers include various organic or inorganic carrier substances which are commonly used as materials for pharmaceutical preparations, such as excipients, lubricants, binders, and disintegrators in solid
20 preparations; solvents, solubilizing agents, suspending agents, isotonizing agents, buffering agents, soothing agents, in liquid preparations. Also, in the pharmaceutical manufacturing process, additives such as antiseptics, antioxidants, coloring agents, sweeteners,
25 absorbents, moistening agents, can be used, if necessary.

Examples of the excipients include lactose, sucrose, D-mannitol, starch, cornstarch, crystalline cellulose, light anhydrous silicic acid.

30 Examples of the lubricants include magnesium stearate, calcium stearate, talc, colloidal silica.

 Examples of the binders include crystalline cellulose, sucrose, D-mannitol, dextrin, hydroxypropylcellulose, hydroxypropylmethylcellulose, polyvinylpyrrolidone, starch, saccharose, gelatin,
35 methylcellulose, carboxymethylcellulose sodium.

 Examples of the disintegrators include starch,

carboxymethylcellulose, carboxymethylcellulose calcium, crosscarmellose sodium, carboxymethylstarch sodium, low-substituted hydroxypropylcellulose (L-HPC).

5 Examples of the solvents include distilled water for injection, alcohol, propylene glycol, macrogol, sesame oil, corn oil.

10 Examples of the solubilizing agents include polyethylene glycol, propylene glycol, D-mannitol, benzyl benzoate, ethanol, trisaminomethane, cholesterol, triethanolamine, sodium carbonate, sodium citrate.

15 Examples of the suspending agents include surfactants such as stearyltriethanolamine, sodium lauryl sulfate, lauryl amino propionic acid, lecithin, benzalkonium chloride, benzethonium chloride, glycerylmonostearate; or hydrophilic polymers such as polyvinyl alcohol, polyvinylpyrrolidone, carboxymethylcellulose sodium, methylcellulose, hydroxymethylcellulose, hydroxyethylcellulose, hydroxypropylcellulose.

20 Examples of the isotonizing agents include glucose, D-sorbitol, sodium chloride, glycerin, D-mannitol.

Examples of the buffering agents include buffer solutions of phosphate, acetate, carbonate and citrate.

Examples of the soothing agents include benzyl alcohol.

25 Examples of the antiseptics include paraoxybenzoates, chlorobutanol, benzyl alcohol, phenethylalcohol, dehydroacetic acid, and sorbic acid.

Examples of the antioxidants include sulfite, ascorbic acid.

30 A MCH antagonist and a pharmaceutical composition of the invention can be safely administered orally or parenterally (e.g. by local, rectal and intravenous administration) in various dosage forms, for instance, as oral drugs such as tablets (including sugar-coated tablets and film-coated tablets), powders, granules, capsules
35 (including soft capsules), solutions; and parenteral

preparations such as injections (e.g. subcutaneous injections, intravenous injections, intramuscular injections, intraperitoneal injections, etc.), external preparations (e.g. nasal preparations, percutaneous preparations, ointments, etc.), suppositories (e.g. rectal suppositories, vaginal suppositories, etc.), sustained-release preparations (e.g. sustained-release microcapsules, etc.), pellets, drip infusions, etc.

The content of compound (I) in a MCH antagonist of the invention and the content of compound (I') in a pharmaceutical composition of the invention are, for instance, about 0.1 to 100 weight percent of the MCH antagonist or whole pharmaceutical composition, respectively.

The dose of a MCH antagonist and a pharmaceutical composition of the invention can be appropriately selected depending on the subject of administration, route of administration, disease, etc.

For instance, the dose per day when a MCH antagonist or a pharmaceutical composition of the invention is orally administered to an adult obesity patient (body weight: about 60 kg), is about 0.1 to about 500 mg, preferably about 1 to about 100 mg, more preferably about 5 to about 100 mg, in terms of compound (I) or compound (I'), each of which is an active ingredient. These amounts can be divided into one to several doses per day for administration.

The MCH antagonist and pharmaceutical composition of the invention can be used in combination with other concomitant drugs which do not interfere with the MCH antagonist and pharmaceutical composition of the invention, for the purpose of "strengthening of therapeutic effect against obesity", "reduction of dose of MCH antagonist", etc. Examples of the concomitant drugs include a "agents for treating diabetes", "agents for treating diabetic complications", "agents for treating obesity other than MCH antagonists", "agents for treating

hypertension", "agents for treating hyperlipidemia (agents for treating arteriosclerosis)", "agents for treating arthritis", "antianxiety agents", "antidepressant". Two or more kinds of these concomitant drugs can be combined
5 in an appropriate ratio for use.

Examples of the above "agents for treating diabetes" include insulin sensitizers, insulin secretion enhancers, biguanides, insulins, α -glucosidase inhibitors, β 3 adrenaline receptor agonists.

10 Examples of the insulin sensitizers include pioglitazone or its salt (preferably hydrochloride), troglitazone, rosiglitazone or its salt (preferably maleate), JTT-501, GI-262570, MCC-555, YM-440, DRF-2593, BM-13-1258, KRP-297, R-119702.

15 Examples of the insulin secretion enhancers include sulfonylureas. Concrete examples of the sulfonylureas include tolbutamide, chlorpropamide, trazamide, acetohexamide, glyclopyramide and its ammonium salt, glibenclamide, gliclazide, glimepiride.

20 Other than the above, examples of insulin secretion enhancers include repaglinide, nateglinide, mitiglinide (KAD-1229), JTT-608.

Examples of biguanides include metformin, buformin, phenformin.

25 Examples of insulins include animal insulins extracted from bovine or porcine pancreas; semi-synthetic human insulin which is enzymatically synthesized from insulin extracted from porcine pancreas; human insulin synthesized by genetic engineering, using Escherichia Coli and yeast. As insulin, also employed are insulin-zinc
30 containing 0.45 to 0.9 (w/w)% of zinc; protamine-insulin-zinc produced from zinc chloride, protamine sulfate and insulin. In addition, insulin can be an insulin fragment or derivative (e.g. INS-1, etc.).

35 Insulin can also include various types such as ultra immediate action type, immediate action type, two-phase

type, intermediate type, prolonged action type, etc., and these can be selected depending on the pathological conditions of patients.

5 Examples of α -glucosidase inhibitors include acarbose, voglibose, miglitol, emiglitate.

Examples of β 3 adrenaline receptor agonists include AJ-9677, BMS-196085, SB-226552, AZ40140.

10 Other than the above, examples of the "agents for treating diabetes" include ergoset, pramlintide, leptin, BAY-27-9955.

Examples of the above "agents for treating diabetic complications" include aldose reductase inhibitors, glycation inhibitors, protein kinase C inhibitors.

15 Examples of aldose reductase inhibitors include torulestat; eparlestat; imirestat; zenarestat; SNK-860; zopolrestat; ARI-509; AS-3201.

Examples of glycation inhibitors include pimagedine.

Examples of protein kinase C inhibitors include NGF, LY-333531.

20 Other than the above, examples of "agents for treating diabetic complications" include alprostadil, thiapride hydrochloride, cilostazol, mexiletine hydrochloride, ethyl eicosapentate, memantine, pimagedline (ALT-711).

25 Examples of the above "agents for treating obesity other than MCH antagonists" include lipase inhibitors and anorectics.

Examples of lipase inhibitors include orlistat.

30 Examples of anorectics include mazindol, dexfenfluramine, fluoxetine, sibutramine, baiamine, (S)-sibutramine, SR-141716, NGD-95-1.

Other than the above, examples of "agents for treating obesity other than MCH antagonists" include lipstatin.

35 Examples of the above "agents for treating hypertension" include angiotensin converting enzyme inhibitors, calcium antagonists, potassium channel openers, angiotensin II antagonists.

Examples of angiotensin converting enzyme inhibitors include captopril, enalapril, alacepril, delapril (hydrochloride), lisinopril, imidapril, benazepril, cilazapril, temocapril, trandolapril, manidipine
5 (hydrochloride).

Examples of calcium antagonists include nifedipine, amlodipine, efonidipine, nicardipine.

Examples of potassium channel openers include levcromakalim, L-27152, AL0671, NIP-121.

10 Examples of angiotensin II antagonists include losartan, candesartan cilexetil, valsartan, irbesartan, CS-866, E4177.

Examples of the above "agents for treating hyperlipidemia (agents for treating arteriosclerosis)"
15 include HMG-CoA reductase inhibitors, fibrate compounds.

Examples of HMG-CoA reductase inhibitors include pravastatin, simvastatin, lovastatin, atorvastatin, fluvastatin, lipantil, cerivastatin, itavastatin, ZD-4522, or their salts (e.g. sodium salts, etc.).

20 Examples of fibrate compounds include bezafibrate, clinofibrate, clofibrate, simfibrate.

Examples of the above "agents for treating arthritis" include ibuprofen.

25 Examples of the above "antianxiety agents" include chlordiazepoxide, diazepam, oxazolam, medazepam, cloxazolam, bromazepam, lorazepam, alprazolam, fludiazepam.

30 Examples of the above "antidepressants" include fluoxetine, fluvoxamine, imipramine, paroxetine, sertraline.

The timing of administration of the above concomitant drugs is not limited. The MCH antagonist or pharmaceutical composition and the concomitant drugs can be administered to the subject simultaneously or at staggered times.

35 The dosages of the concomitant drugs can be determined in accordance with clinically used dosages, and can be

appropriately selected according to the subject of administration, route of administration, diseases and combinations of drugs, etc.

5 The administration forms for the concomitant drugs are not particularly limited as long as a MCH antagonist or a pharmaceutical composition are used in combination with a concomitant drugs at the time of administration. Examples of such administration forms includes 1) administration of a single preparation obtained by simultaneous preparation
10 of MCH antagonist or pharmaceutical composition together with concomitant drugs, 2) simultaneous administration of two kinds of preparations obtained by separate preparation of MCH antagonist or pharmaceutical composition, and concomitant drugs, through the same route of
15 administration, 3) staggered administration of two kinds of preparations obtained by separate preparation of MCH antagonist or pharmaceutical composition, and concomitant drugs, through the same route of administration, 4) simultaneous administration of two kinds of preparations
20 obtained by separate preparation of MCH antagonist or pharmaceutical composition, and concomitant drugs, through different routes of administration, 5) staggered administration of two kinds of preparations obtained by separate preparation of MCH antagonist or pharmaceutical
25 composition, and concomitant drugs, through different routes of administration (for instance, administration of MCH antagonist or pharmaceutical composition; and concomitant drugs in this order; or administration in reverse order).

30 The ratio of combination of MCH antagonist or pharmaceutical composition with concomitant drugs can be appropriately selected in accordance with the subject of administration, route of administration and diseases, etc.

35 This invention further relates to "a pharmaceutical comprising a melanin-concentrating hormone antagonist in

combination with at least one species selected from the group consisting of an agent for treating diabetes, an agent for treating hypertension and an agent for treating arteriosclerosis".

5 Here, the "melanin-concentrating hormone antagonist" is not especially limited as long as it is a compound having a melanin-concentrating hormone antagonistic action, and may be either of a peptide compound or a non-peptide compound.

10 As "an agent for treating diabetes", "an agent for treating hypertension" and "an agent for treating arteriosclerosis", those exemplified as the above concomitant drugs can be mentioned.

15 These drugs can be used in the same manner as in the above "combination of MCH antagonist of the invention with concomitant drugs".

20 The pharmaceutical provides excellent effects such as "strengthening of therapeutic effect against obesity", "reduction of dose of MCH antagonist", etc. as compared to single use of each drug.

BEST MODE FOR CARRYING OUT THE INVENTION

25 This invention will be explained further in detail by the following Reference Examples, Examples, Preparation Examples, and Experimental Examples. However, these do not limit this invention, and they can be changed within the scope that does not deviate from the scope of this invention.

30 In the following Reference Examples and Examples, "room temperature" means 0 to 30°C. Anhydrous magnesium sulfate or anhydrous sodium sulfate was used to dry the organic layer. "%" means percent by weight, unless otherwise specified.

35 Infrared absorption spectra were determined by the diffuse reflectance method, using fourier transform type infrared spectrophotometer.

FABMS (pos) is mass spectrum determined by the (+) method, in Fast Atom Bombardment Mass Spectrometry.

Other symbols used in the description have the following meanings.

	s	:	singlet
	d	:	doublet
	t	:	triplet
	q	:	quartet
10	m	:	multiplet
	br	:	broad
	J	:	coupling constant
	Hz	:	Hertz
	CDCl ₃	:	heavy chloroform
15	DMSO-d ₆	:	heavy dimethylsulfoxide
	THF	:	tetrahydrofuran
	DMF	:	N,N-dimethylformamide
	DMSO	:	dimethylsulfoxide
	WSCD	:	1-ethyl-3-(3-dimethylaminopropyl)
20		:	carbodiimide
	WSC	:	1-ethyl-3-(3-dimethylaminopropyl)
		:	carbodiimide hydrochloride
	¹ H-NMR	:	proton nuclear resonance
		:	(Free substances were usually measured in
25		:	CDCl ₃ .)
	IR	:	infrared absorption spectrum
	Me	:	methyl
	Et	:	ethyl
	HOBt	:	1-hydroxy-1H-benzotriazole
30	IPE	:	diisopropyl ether
	DMAP	:	4-dimethylaminopyridine

In this specification and drawings, when bases and amino acids are shown by codes, these codes are based on those by the IUPAC-IUB Commission on Biochemical

Nomenclature or common codes in the concerned fields.
Examples of these codes are shown below. Also, where some optical isomers of amino acids can exist, the L form is shown unless otherwise specified.

5	DNA	: deoxyribonucleic acid
	cdNA	: complementary deoxyribonucleic acid
	A	: adenine
	T	: thymine
	G	: guanine
10	C	: cytosine
	RNA	: ribonucleic acid
	mRNA	: messenger ribonucleic acid
	dATP	: deoxyadenosine triphosphate
	dTTP	: deoxythymidine triphosphate
15	dGTP	: deoxyguanosine triphosphate
	dCTP	: deoxycytidine triphosphate
	ATP	: adenosine triphosphate
	EDTA	: ethylenediamine tetraacetic acid
	SDS	: sodium dodecyl sulfate
20	EIA	: enzyme immunoassay
	Gly	: glycine
	Ala	: alanine
	Val	: valine
	Leu	: leucine
25	Ile	: isoleucine
	Ser	: serine
	Thr	: threonine
	Cys	: cysteine
	Met	: methionine
30	Glu	: glutamic acid
	Asp	: aspartic acid
	Lys	: lysine
	Arg	: arginine
	His	: histidine
35	Phe	: phenylalanine
	Tyr	: tyrosine

	Tro	: tryptophan
	Pro	: proline
	Asn	: asparagine
	Gln	: glutamine
5	pGl	: pyroglutamine
	Me	: methyl group
	Et	: ethyl group
	Bu	: butyl group
	Ph	: phenyl group
10	TC	: thiazolidine-4(R)-carboxamide group

Substituents, protecting groups and reagents frequently used in this specification, are shown by the following symbols.

15	Tos	: p-toluenesulfonyl
	CHO	: formyl
	Bzl	: benzyl
	Cl ₂ Bzl	: 2,6-dichlorobenzyl
	Bom	: benzyloxymethyl
20	Z	: benzyloxycarbonyl
	Cl-Z	: 2-chlorobenzyloxycarbonyl
	Br-Z	: 2-bromobenzyloxycarbonyl
	Boc	: t-butoxycarbonyl
	DNP	: dinitrophenol
25	Trt	: trityl
	Bum	: t-butoxymethyl
	Fmoc	: N-9-fluorenylmethoxycarbonyl
	HOBt	: 1-hydroxybenztriazole
	HOObt	: 3,4-dihydro-3-hydroxy-4-oxo-1,2,3-
30		benzotriazine
	HONB	: 1-hydroxy-5-norbornene-2,3-
		dicarbodiimide
	DCC	: N,N'-dicyclohexylcarbodiimide

35 SEQ ID NO in the SEQUENCE LISTING in the specification of the present application shows the following sequences.

- [SEQ ID NO : 1] shows a synthetic DNA used for screening of cDNA coding rat SLC-1.
- 5 [SEQ ID NO : 2] shows a synthetic DNA used for screening of cDNA coding rat SLC-1.
- [SEQ ID NO : 3] shows an entire amino acid sequence of rat SLC-1.
- [SEQ ID NO : 4] shows an entire base sequence of rat SLC-1cDNA wherein Sal I recognition sequence was added to the 5' side,
10 and Spe I recognition sequence was added to the 3' side.
- [SEQ ID NO : 5] shows riboprobe used to determine the quantity of SLC-1mRNA expressed in each clone of rat SLC-1 expression CHO cells.
- 15 [SEQ ID NO : 6] shows a synthetic DNA used to obtain cDNA for coding of human SLC-1.
- [SEQ ID NO : 7] shows a primer used to make double-strand cDNA for coding human SLC-1.
- [SEQ ID NO : 8] shows an entire base sequence of cDNA for coding human SLC-1.
- 20 [SEQ ID NO : 9] shows an entire amino acid sequence of human SLC-1.
- [SEQ ID NO : 10] shows a synthetic DNA used for screening of cDNA for coding human SLC-1(S).
- 25 [SEQ ID NO : 11] shows a synthetic DNA used for screening of cDNA for coding human SLC-1(S).
- [SEQ ID NO : 12] shows a synthetic DNA used for screening of cDNA for coding human SLC-1(L).
- 30 [SEQ ID NO : 13] shows a synthetic DNA used for screening of cDNA for coding human SLC-1(L).
- [SEQ ID NO : 14] shows an entire base sequence of human SLC-1(S) cDNA wherein Sal I recognition sequence was added to the 5' side, and Spe I recognition sequence was added to the 3' side.
- 35 [SEQ ID NO : 15] shows an entire base sequence of human SLC-1(L) cDNA wherein Sal I recognition sequence was added

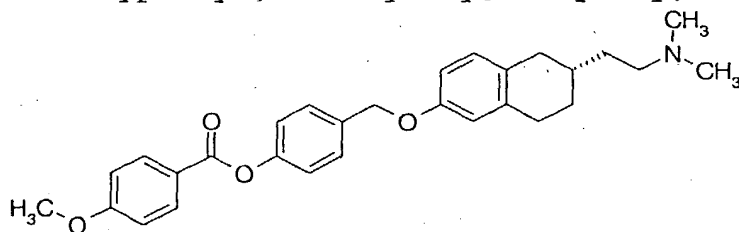
to the 5' side, and Spe I recognition sequence was added to the 3' side.

[SEQ ID NO : 16] shows riboprobe used to determine the quantity of SLC-1mRNA expressed in each clone of human SLC-1(S) expression CHO cells and SLC-1(L) expression CHO cells.

Transformant *Escherichia coli* DH10B/phSLC1L8 transformed by plasmid containing DNA which codes the base sequence shown by SEQ ID NO : 9, obtained in Reference Example 1 - 6, is on deposit with National Institute of Bioscience and Human-Technology (NIBH), Agency of Industrial Science and Technology, Ministry of International Trade and Industry, as deposit number FERM BP-6632 from February 1, 1999; and with the Institute for Fermentation, Osaka, Japan (IFO), as deposit number IFO 16254 from January 21, 1999.

Reference Example 1

2-(R)-[2-(N,N-Dimethylamino)ethyl]-6-(4-[(4-methoxyphenyl)carbonyloxy]benzyloxy)tetralin



Diethyl azodicarboxylate (40% toluene solution, 0.95 g) was added dropwise to THF solution (6 ml) of 2-(R)-[2-(N,N-dimethylamino)ethyl]-6-hydroxytetralin (300 mg), 4-(hydroxymethyl)phenyl 4-methoxybenzoate (530 mg), and triphenylphosphine (430 mg) under ice-cooling. After stirring for 2 hours at room temperature, the reaction mixture was concentrated. The residue was purified using alumina column chromatography (development solvent; hexane ~ hexane : ethyl acetate = 10:1), and the titled compound

(320 mg) was obtained after recrystallization (ethyl acetate-hexane).

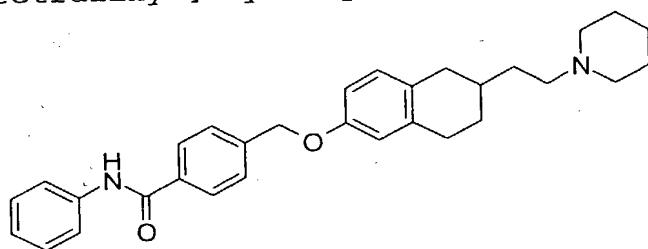
Melting point: 111 - 114°C

$[\alpha]_D^{20} = +44.4^\circ$ (c = 0.502, methanol)

5

Reference Example 2

N-Phenyl-4-[[2-(2-piperidinoethyl)-6-tetralinyl]oxymethyl]benzamide



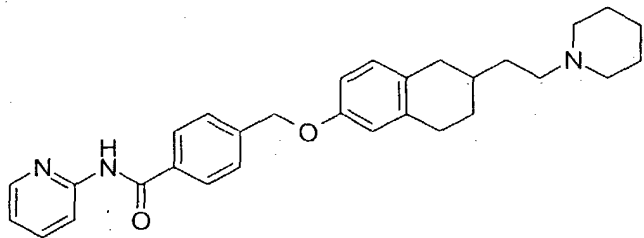
10 Triethylamine (0.11 ml) was added to THF suspension (3 ml) of 4-[[2-(2-piperidinoethyl)-6-tetralinyl]oxymethyl]benzoate (300 mg). Further, THF solution (0.5 ml) of trimethylacetyl chloride (92 mg) was added dropwise under ice-cooling, which was stirred for 30 minutes. The temperature of the reaction mixture was raised to room temperature, which was stirred for 1 hour.

15 THF solution (0.5 ml) of aniline (85 mg) was added dropwise to the reaction mixture under ice-cooling, which was stirred for 1 hour. After the reaction mixture was stirred for 24 hours at room temperature, saturated sodium bicarbonate solution was added, and extraction was conducted using a mixed solution of ethyl acetate and THF.

20 The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was recrystallized from THF-methanol-IPE to give the titled compound (150 mg).
25 Melting point: 183 - 185°C

Reference Example 3

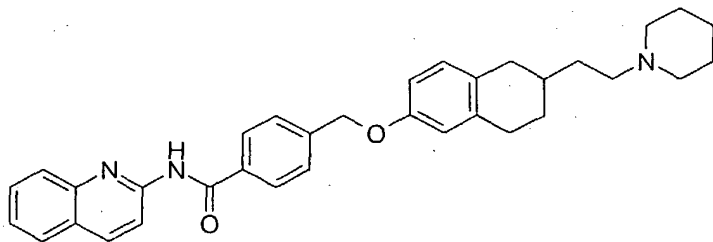
30 4-[[2-(2-Piperidinoethyl)-6-tetralinyl]oxymethyl]-N-(2-pyridinyl)benzamide



Triethylamine (0.11 ml) was added to THF suspension (6 ml) of 4-[[2-(2-piperidinoethyl)-6-tetralinyl]oxymethyl]benzoate (300 mg). Trimethylacetyl chloride (0.095 ml) was added dropwise to the obtained suspension under ice-cooling, which was stirred for 30 minutes. The temperature of the reaction mixture was raised to room temperature, which was stirred for 1 hour. THF solution (1.0 ml) of 2-aminopyridine (110 mg) was added dropwise to the reaction mixture under ice-cooling, which was stirred for 1 hour. Then the reaction mixture was stirred at room temperature for 6 hours, and at 60°C for 12 hours, which was refluxed with heating for 6 hours. Saturated sodium bicarbonate solution was added to the reaction mixture, and extraction was conducted using ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was purified using alumina column chromatography (development solvent: THF), and recrystallized (ethyl acetate-IPE) to give the titled compound (30 mg).

Reference Example 4

4-[[2-(2-Piperidinoethyl)-6-tetralinyl]oxymethyl]-N-(2-quinolinyl)benzamide

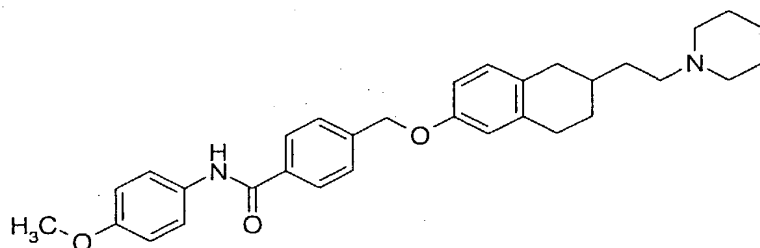


Triethylamine (0.22 ml) was added to THF suspension (6 ml) of 4-[[2-(2-piperidinoethyl)-6-tetralinyl]oxymethyl]benzoate (300 mg). Further, trimethylacetyl chloride (0.095 ml) was added dropwise to under ice-cooling, which was stirred for 30 minutes. The temperature of the reaction mixture was raised to room temperature, which was stirred for 1 hour. THF solution (1.0 ml) of 2-aminoquinoline (170 mg) was added dropwise to the reaction mixture under ice-cooling, which was stirred at room temperature for 12 hours. Saturated sodium bicarbonate solution was added to the reaction mixture, and extraction was conducted using ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was purified using alumina column chromatography (development solvent: THF), and recrystallized (ethyl acetate-diisopropyl ether) to give the titled compound (45 mg).

Melting point: 135 - 138° C

Reference Example 5

N-(4-Methoxyphenyl)-4-[[2-(2-piperidinoethyl)-6-tetralinyl]oxymethyl]benzamide



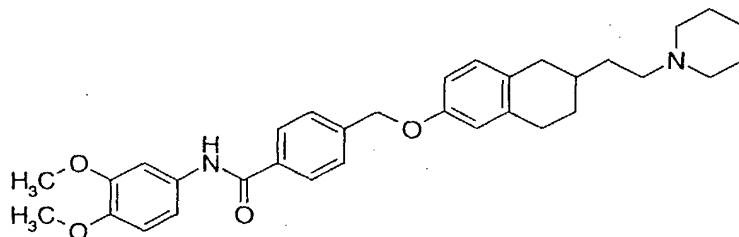
WSCD (0.11 ml) was added to DMF solution (2 ml) of 4-[[2-(2-piperidinoethyl)-6-tetralinyl]oxymethyl]benzoate (170 mg), 4-methoxyaniline (53 mg), HOBt (70 mg) and DMAP (60 mg) at room temperature, which was stirred for 12 hours. 10% aqueous potassium carbonate solution and water was added to the reaction mixture, and extraction was conducted using a mixed

solution of THF and ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was purified using alumina column chromatography (development solvent: THF), and recrystallized (THF-IPE) to give the

5 titled compound (140 mg).
Melting point: 193 - 196°C

Reference Example 6

10 N-(3,4-Dimethoxyphenyl)-4-[[2-(2-piperidinoethyl)-6-tetralinyl]oxymethyl]benzamide

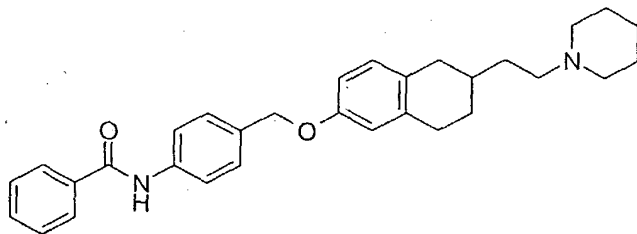


WSCD (free form, 0.2 ml) was added to DMF solution (3 ml) of 4-[[2-(2-piperidinoethyl)-6-tetralinyl]oxymethyl]benzoate (300 mg), 3,4-dimethoxyaniline (120 mg), HOBt (120 mg) and DMAP (100 mg) at room temperature, which was stirred for 12 hours. 10% aqueous potassium carbonate solution was added to the reaction mixture, and the resulting crystals were collected by filtration. The crystals were washed with water, then dried. The crystals were purified using alumina column chromatography (development solvent: THF), and recrystallized (THF-IPE) to give the titled compound (330 mg).

20
25 Melting point: 178 - 180°C

Reference Example 7

6-[4-(Benzoylamino)benzyloxy]-2-(2-piperidinoethyl)tetralin



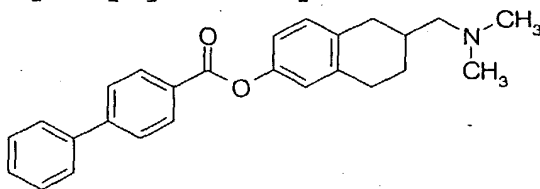
Sodium hydride (60% oily, 85 mg) was added to DMF solution of 6-hydroxy-2-(2-piperidinoethyl)tetralin (500 mg) at room temperature, which was stirred for 1 hour.

- 5 N-[4-(bromomethyl)phenyl]benzamide (670 mg) was added to the reaction mixture at room temperature, which was stirred for 1 hour. Water was added to the reaction mixture, and extraction was conducted using ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was purified using alumina column chromatography (development solvent : THF), and recrystallized (ethyl acetate) to give the titled compound (200 mg).
10 Melting point: 176 - 179° C

15

Reference Example 8

2-[(N,N-Dimethylamino)methyl]-6-tetralinyl 4-biphenylcarboxylate



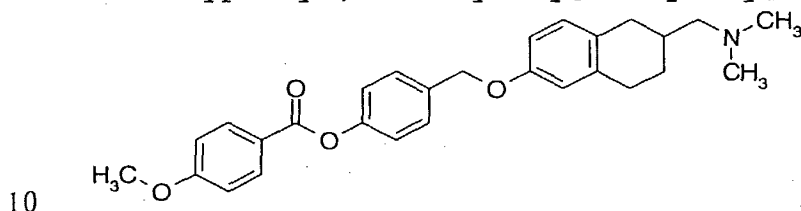
- 20 4-Biphenylcarboxylic acid (580 mg) and WSC (560 mg) were added to pyridine solution (6 ml) of 2-[(N,N-dimethylamino)methyl]-6-hydroxytetralin (300 mg), which was stirred at room temperature for 36 hours. Saturated sodium bicarbonate solution and water were added to the
25 reaction mixture, and extraction was conducted using ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then

concentrated. The residue was purified using alumina column chromatography (development solvent; hexane ~ hexane:ethyl acetate = 10:1), and recrystallized (hexane) to give the titled compound (300 mg).

5 Melting point: 85 - 86°C

Reference Example 9

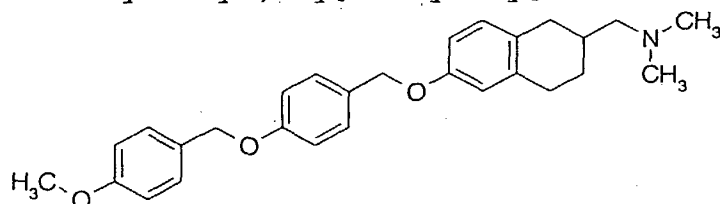
2-[(N,N-Dimethylamino)methyl]-6-[4-[(4-methoxyphenyl)carbonyloxy]benzyloxy]tetralin



Diethyl azodicarboxylate (40% toluene solution, 950 mg) was added dropwise to THF solution (3 ml) of 2-[(N,N-dimethylamino)methyl]-6-hydroxytetralin (150 mg), 4-(hydroxymethyl)phenyl 4-methoxybenzoate (570 mg) and triphenylphosphine (574 mg) at room temperature, which was stirred for 3 hours. The reaction mixture was concentrated, and the residue was purified using alumina column chromatography (development solvent; hexane ~ hexane:ethyl acetate = 6:1), and recrystallized (ethyl acetate-hexane) to give the titled compound (175 mg).
 15
 20 Melting point: 119 - 121°C

Reference Example 10

25 2-[(N,N-Dimethylamino)methyl]-6-[4-[(4-methoxybenzyl)oxy]benzyloxy]tetralin



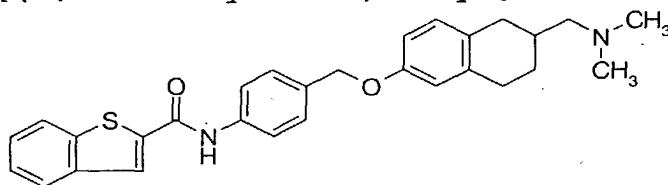
Diethyl azodicarboxylate (40% toluene solution, 1.91 g) was added dropwise to THF solution (6 ml) of 2-

[(N,N-dimethylamino)methyl]-6-hydroxytetralin (300 mg),
4-[(4-methoxybenzyl)oxy]benzylalcohol (1.07 g) and
triphenylphosphine (1.15g) at room temperature, which was
stirred for 12 hours. The reaction mixture was
5 concentrated, and the residue was purified using alumina
column chromatography (development solvent; hexane -
hexane:ethyl acetate = 10:1), and recrystallized (ethyl
acetate-hexane) to give the titled compound (260 mg).
Melting point: 106 - 111°C

10

Reference Example 11

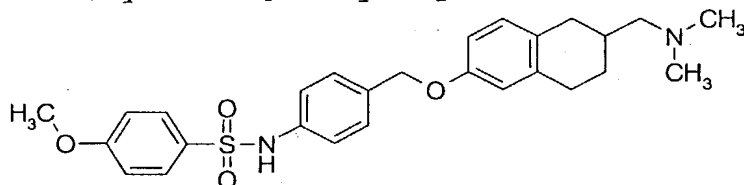
6-[4-[(1-Benzothiophen-2-yl)carbonylamino]benzyloxy]-2-
[(N,N-dimethylamino)methyl]tetralin



15 One drop of DMF was added to THF solution (4 ml) of
1-benzothiophene-2-carboxylic acid (230 mg), and oxalyl
chloride (0.23 ml) was further added under ice-cooling,
which was stirred for 30 minutes at room temperature. The
reaction mixture was concentrated, which was dissolved in
20 THF (1 ml). The obtained solution was added dropwise to
pyridine solution (6 ml) of 6-(4-aminobenzyl)-2-
[(N,N-dimethylamino)methyl]tetralin (300 mg), which was
stirred for 15 minutes. After stirring at room temperature
for another 15 minutes, 10% aqueous potassium carbonate
25 solution was added to the reaction mixture, and extraction
was conducted using ethyl acetate. The organic layer was
washed with water and saturated aqueous sodium chloride
solution, dried, and then concentrated. The residue was
purified using alumina column chromatography (development
30 solvent; ethyl acetate), and recrystallized (THF-IPE) to
give the titled compound (250 mg).
Melting point: 165 - 169°C

Reference Example 12

2-[(N,N-Dimethylamino)methyl]-6-[4-[(4-methoxyphenyl)sulfonylamino]benzyloxy]tetralin



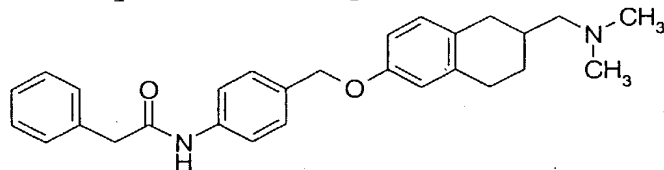
5

THF solution (1 ml) of 4-methoxybenzenesulfonyl chloride (270 mg) was added dropwise to pyridine solution (6 ml) of 6-[(4-aminobenzyl)oxy]-2-[(N,N-dimethylamino)methyl]tetralin (300 mg) under ice-cooling, which was stirred for 15 minutes. After stirring at room temperature for further 15 minutes, 10% aqueous potassium carbonate solution was added to the reaction mixture, and extraction was conducted using ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was purified using alumina column chromatography (development solvent: ethyl acetate), and recrystallized (ethyl acetate-IPE) to give the titled compound (260 mg). Melting point: 137 - 140°C

20

Reference Example 13

6-[4-(Benzylcarbonylamino)benzyloxy]-2-[(N,N-dimethylamino)methyl]tetralin



25

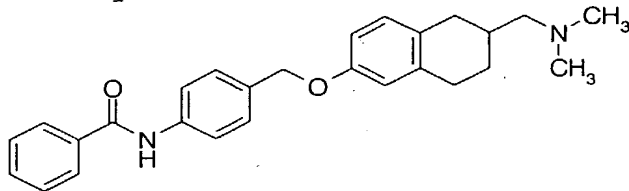
THF solution (1 ml) of phenylacetyl chloride (200 mg) was added dropwise to pyridine solution (6 ml) of 6-[(4-aminobenzyl)oxy]-2-[(N,N-dimethylamino)methyl]tetralin (300 mg) under ice-cooling, which was stirred for 15 minutes. After stirring at room temperature for further 15 minutes, saturated sodium

30

bicarbonate solution was added to the reaction mixture, and extraction was conducted using ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was purified using alumina column chromatography (development solvent; hexane ~ hexane:ethyl acetate = 2:1), and recrystallized to give the titled compound (175 mg). Melting point: 130 - 135°C

10 Reference Example 14

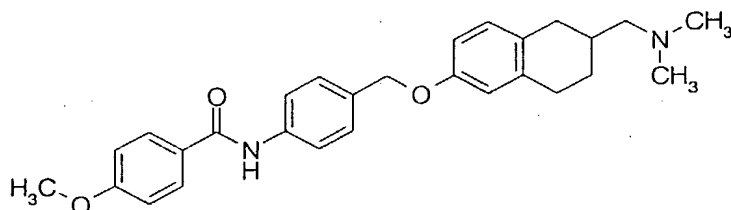
6-[4-(Benzoylamino)benzyloxy]-2-[(N,N-dimethylamino)methyl] tetralin



Benzoyl chloride (0.14 ml) was added dropwise to pyridine solution (6 ml) of 6-[(4-aminobenzyl)oxy]-2-[(N,N-dimethylamino)methyl]tetralin (300 mg) under ice-cooling, which was stirred at room temperature for 30 minutes. 10% aqueous potassium carbonate solution was added to the reaction mixture, and extraction was conducted using ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was purified using alumina column chromatography (development solvent; ethyl acetate), and recrystallized (THF-IPE) to give the titled compound (240 mg). Melting point: 128 - 133°C

Reference Example 15

2-[(N,N-Dimethylamino)methyl]-6-[4-[(4-methoxybenzoyl)amino]benzyloxy]tetralin

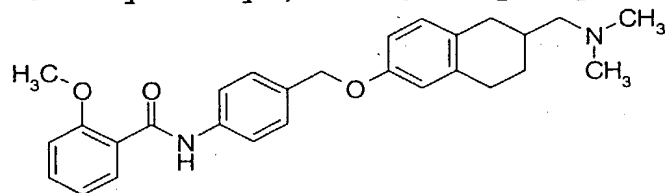


p-Anisoyl chloride (0.20 ml) was added dropwise to pyridine solution (6 ml) of 6-[(4-aminobenzyl)oxy]-2-[(N,N-dimethylamino)methyl]tetralin (300 mg) under ice-cooling, which was stirred at room temperature for 30 minutes. 10% aqueous potassium carbonate solution was added to the reaction mixture, and extraction was conducted using ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was purified using alumina column chromatography (development solvent: ethyl acetate), and recrystallized (THF-IPE) to give the titled compound (300 mg).

Melting point: 155 - 159° C

Reference Example 16

2-[(N,N-Dimethylamino)methyl]-6-[4-[(2-methoxybenzoyl)amino]benzyloxy]tetralin



o-Anisoyl chloride (0.15 ml) was added dropwise to pyridine solution (4 ml) of 6-[(4-aminobenzyl)oxy]-2-[(N,N-dimethylamino)methyl]tetralin (200 mg) under ice-cooling, which was stirred at room temperature for 30 minutes. 10% aqueous potassium carbonate solution was added to the reaction mixture, and extraction was conducted using ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was purified

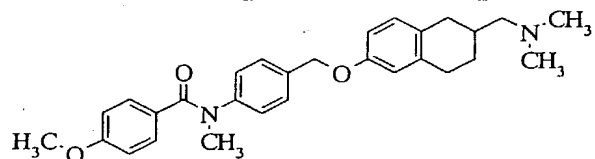
using alumina column chromatography (development solvent; THF), and recrystallized (ethyl acetate-hexane) to give the titled compound (200 mg).

Melting point: 106 - 108°C

5

Reference Example 17

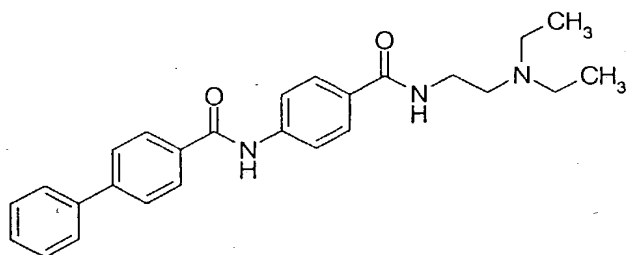
6-[4-[N-(4-Methoxybenzoyl)-N-methylamino]benzyloxy]-2-[(N,N-dimethylamino)methyl]tetralin



- 10 Diethyl azodicarboxylate (40% toluene solution, 960 mg) was added dropwise to THF solution (3 ml) of 2-[(N,N-dimethylamino)methyl]-6-hydroxytetralin (150 mg),
- 15 N-[4-(hydroxymethylphenyl)-4-methoxy-N-methylbenzamide (600 mg) and triphenylphosphine (570 mg) at room temperature, which was stirred for 12 hours. After the reaction mixture was concentrated, the residue was purified using silica gel column chromatography (development solvent; hexane ~ ethyl acetate ~ ethyl acetate:methanol = 1:2), and then purified using alumina column
- 20 chromatography (development solvent; hexane ~ hexane:ethyl acetate = 2:1) to give the titled compound (185 mg). ¹H-NMR (CDCl₃) δ: 1.20-1.50(1H, m), 1.80-2.46(5H, m), 2.25(6H, s), 2.68-2.86(3H, m), 3.47(3H, s), 3.74(3H, s), 4.95(2H, s), 6.52-6.76(4H, m), 6.84-7.14(3H, m), 7.22-7.38(4H, m).
- 25

Reference Example 18

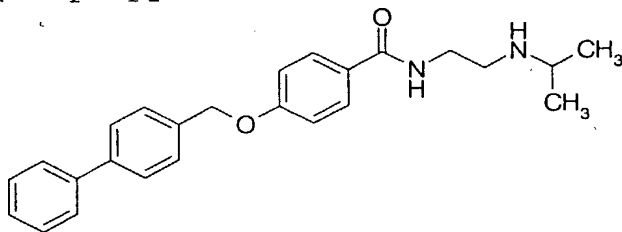
N-[4-[[[2-(Diethylamino)ethyl]amino]carbonyl]phenyl] 4-biphenyllylcarboxamide



Oxalyl chloride (0.46 ml) and DMF (1 drop) were added to THF solution (15 ml) of 4-biphenylcarboxylic acid (0.879g) under ice-cooling. The reaction mixture was stirred at room temperature for 30 minutes, and concentrated. The residue was dissolved in THF (10 ml), which was added dropwise to THF (20 ml) suspension of procaineamide hydrochloride (1.078 g) and triethylamine (1.4 ml) at 0°C. After stirring at 0°C for 30 minutes, 10% aqueous potassium carbonate solution was added to the reaction mixture, and extraction was conducted using ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was recrystallized using methanol to give the titled compound (1.147 g). Melting point: 237 - 240°C (decomposition)

Reference Example 19

4-(4-Biphenylmethoxy)-N-[2-(isopropylamino)ethyl]benzamide



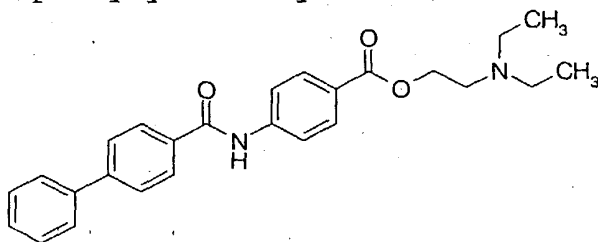
WSC (0.708 g), HOBt (0.521 g), N-isopropyl ethylenediamine (0.353 g) and triethylamine (1 ml) were added to a mixed solution of 4-(4-biphenylmethoxy) benzoate (1.007 g) in THF (30 ml) and acetonitrile (30 ml). After stirring at room temperature for 18 hours, water was added to the reaction mixture, and extraction was conducted

using ethyl acetate. The organic layer was washed with 10% aqueous potassium carbonate solution and saturated aqueous sodium chloride solution, dried, and then concentrated.

The residue was recrystallized using ethanol to give the
5 titled compound (0.806 g).
Melting point: 150 - 154°C

Reference Example 20

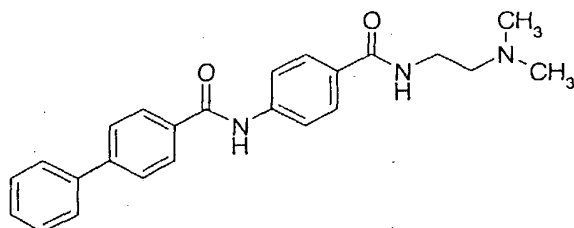
2-(N,N-Diethylamino)ethyl 4-(4-
10 biphenylcarboxylamino)benzoate



Oxalyl chloride (0.39 ml) and DMF (1 drop) were added to THF solution (15 ml) of 4-biphenylcarboxylic acid (1.091 g) under ice-cooling, which was stirred at room
15 temperature for 30 minutes, and concentrated. The residue was dissolved in THF (10 ml), which was added dropwise to THF suspension (30 ml) of procaine hydrochloride (1.091 g) and triethylamine (0.67 ml) at 0°C. After stirring at 0°C
20 for 30 minutes, 10% aqueous potassium carbonate was added to the reaction mixture, and extraction was conducted using ethyl acetate. The organic layer was washed with saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was recrystallized using ethyl acetate/hexane to give the titled compound (0.728 g).
25 Melting point: 146 - 149°C

Reference Example 21

N-[4-[[[2-(Dimethylamino)ethyl]amino]carbonyl]phenyl]
4-biphenylcarboxamide



WSC (0.248 g), HOBt (0.156 g), N,N-dimethyl ethylenediamine (0.097 g) and triethylamine (0.21 ml) were added to a mixed solution of 4-(4-

5 biphenylcarbonylamino)benzoate (0.323 g) in THF (15 ml) and acetonitrile (15 ml). After stirring at room temperature for 18 hours, water was added to the reaction mixture, and extraction was conducted using ethyl acetate.

The organic layer was washed with 10% aqueous potassium carbonate and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was recrystallized using methanol/diethyl ether to give the

10 titled compound (0.100 g).
Melting point: 261 - 264°C (decomposition)

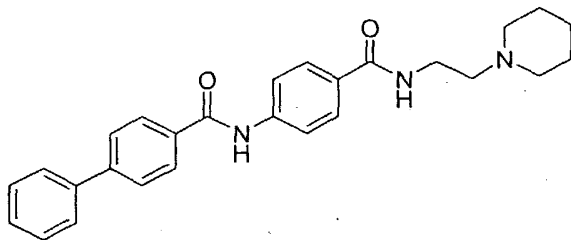
15

The compounds described in the following Reference Examples 22 to 25 were produced in the same manner as in Reference Example 21.

20

Reference Example 22

N-[4-[[2-(Piperidinoethyl)amino]carbonyl]phenyl] 4-biphenylcarboxamide

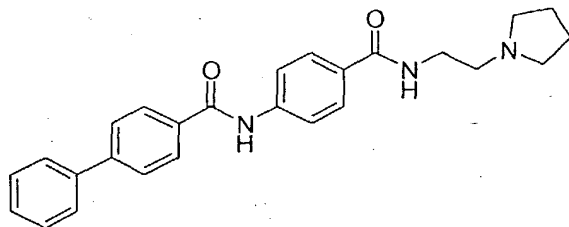


Melting point: 247 - 252°C (decomposition)

25

Reference Example 23

N-[4-[[2-(1-Pyrrolidinyl)ethyl]amino]carbonyl]phenyl]
4-biphenyllylcarboxamide

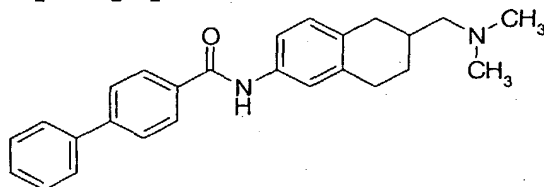


Melting point: 241 - 245°C (decomposition)

5

Reference Example 24

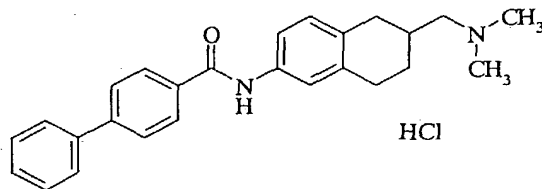
N-[2-(N,N-Dimethylamino)methyl-6-tetralinyl]-4-
biphenyllylcarboxamide



10 Melting point: 164 - 166°C

Reference Example 25

N-[2-(N,N-Dimethylamino)methyl-6-tetralinyl]-4-
biphenyllylcarboxamide hydrochloride



HCl

15

Melting point: >250°C

¹H-NMR δ : 1.24-1.54 (1H, m), 1.84-2.10 (2H, m), 2.20-2.50
(3H, m), 2.26 (6H, s), 2.79-3.01 (3H, m), 7.10 (1H, d,
J=8Hz), 7.28-7.54 (5H, m), 7.60-7.82 (5H, m), 7.94 (2H, d,
J=8Hz).

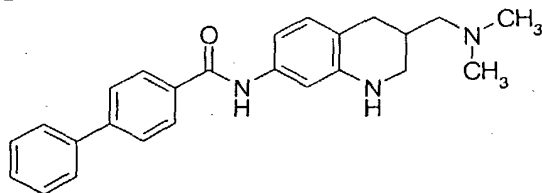
20

IR(KBr) 3028, 2910, 2640, 1658, 1538, 1417, 746, 701 cm⁻¹

Reference Example 26

N-[3-[(N,N-Dimethylamino)methyl]-1,2,3,4-tetrahydro-7-

quinolinyl]-4-biphenylcarboxamide

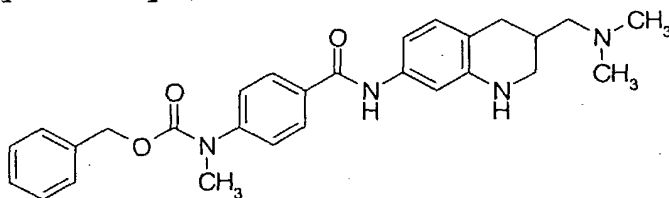


One drop of DMF was added to THF solution of 4-biphenylcarboxylic acid (145 mg), and oxalyl chloride
 5 (0.1 ml) was added dropwise to the solution under ice-cooling, which was stirred at room temperature for 30 minutes. After the reaction mixture was concentrated, the residue was dissolved in THF (1 ml), which was added dropwise to pyridine solution (1.5 ml) of 7-amino-3-
 10 [(N,N-dimethylamino)methyl]-1,2,3,4-tetrahydroquinoline (150 mg) under ice-cooling, and the reaction mixture was stirred for 30 minutes. After the temperature of the reaction mixture was raised to room temperature, 10% aqueous potassium carbonate was added to the reaction
 15 mixture, and extraction was conducted using a mixed solution of THF and ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was recrystallized using THF-IPE to give the titled compound
 20 (180 mg).

Melting point: 206 - 211°C

Reference Example 27

4-[N-[(Benzyloxy)carbonyl]-N-methylamino]-N-[3-[(N,N-
 25 dimethylamino)methyl]-1,2,3,4-tetrahydro-7-quinolinyl]benzamide



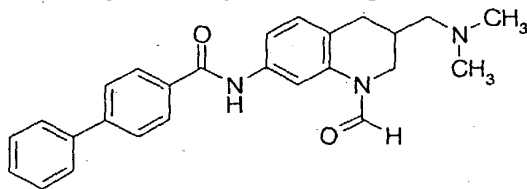
One drop of DMF was added dropwise to THF solution (2

ml) of 4-[N-[(benzyloxy)carbonyl]-N-methylamino]benzoic acid (210 mg), and then oxalyl chloride (0.1 ml) was added dropwise under ice-cooling, which was stirred at room temperature for 30 minutes. After the reaction mixture was concentrated, the residue was dissolved in THF (1 ml), which was added dropwise to pyridine solution (1.5 ml) of 7-amino-3-[(N,N-dimethylamino)methyl]-1,2,3,4-tetrahydroquinoline (150 mg) under ice-cooling. The reaction mixture was then stirred for 30 minutes. After the temperature of the reaction mixture was raised to room temperature, 10% aqueous potassium carbonate solution was added, and extraction was conducted using a mixed solution of THF and ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was recrystallized using THF-IPE to give the titled compound (220 mg).

Melting point: 167 - 172°C

Reference Example 28

N-[3-[(N,N-Dimethylamino)methyl]-1-formyl-1,2,3,4-tetrahydro-7-quinolinyl]-4-biphenylcarboxamide



Anhydrous acetic acid (0.1 ml) was added to formic acid (1 ml), which was stirred at 55°C for 2 hours. N-[3-[(N,N-dimethylamino)methyl]-1,2,3,4-tetrahydro-7-quinolinyl]-4-biphenylcarboxamide (80 mg) was added to the reaction mixture under ice-cooling, which was stirred at room temperature for 72 hours. 10% aqueous potassium carbonate solution was added to the reaction mixture to make the mixture alkaline, and extraction was conducted using ethyl acetate. The organic layer was washed with water and

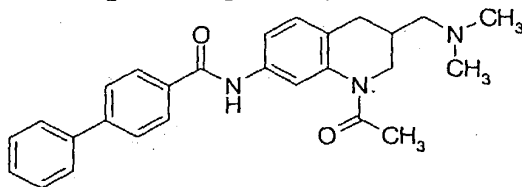
saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was recrystallized using THF-IPE to give the titled compound (80 mg).

Melting point: 134 - 138°C

5

Reference Example 29

N-[1-Acetyl-3-[(N,N-dimethylamino)methyl]-1,2,3,4-tetrahydro-7-quinolyl]-4-biphenylcarboxamide



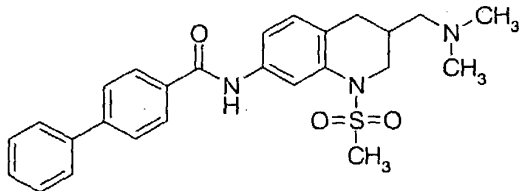
- 10 Acetyl chloride (0.02 ml) was added to pyridine solution (1 ml) of N-[3-[(N,N-dimethylamino)methyl]-1,2,3,4-tetrahydro-7-quinolyl]-4-biphenylcarboxamide (80 mg) under ice-cooling, which was stirred for 15 minutes, and then stirred at room temperature for 15 minutes. 10% aqueous potassium carbonate solution was added to the reaction mixture, and extraction was conducted using ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was recrystallized using THF-IPE to give the titled compound (64 mg).

15

Melting point: 167 - 173°C

Reference Example 30

- 25 N-[3-[(N,N-Dimethylamino)methyl]-1-methylsulfonyl-1,2,3,4-tetrahydro-7-quinolyl]-4-biphenylcarboxamide



Methanesulfonyl chloride (0.02 ml) was added to

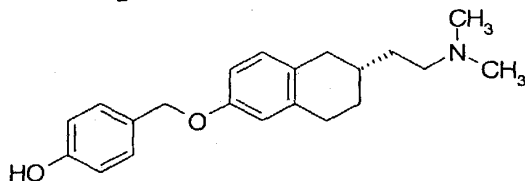
pyridine solution (1 ml) of N-[3-[(N,N-dimethylamino)methyl]-1,2,3,4-tetrahydro-7-quinolinyl]-4-biphenylcarboxamide (80 mg) under ice-cooling, which was stirred at room temperature for 1 hour. Further,
 5 methanesulfonyl chloride (0.02 ml) was added to the reaction mixture under ice-cooling, which was stirred at room temperature for 12 hours. 10% aqueous potassium carbonate solution was added to the reaction mixture, and extraction was conducted using ethyl acetate. The organic
 10 layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was recrystallized using THF-IPE to give the titled compound (64 mg).

Melting point: 184 - 188°C

15

Reference Example 31

2-(R)-[2-(N,N-Dimethylamino)ethyl]-6-(4-hydroxyphenyl) methoxytetralin



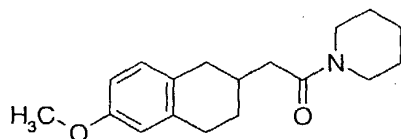
20 THF solution (2 ml) of 2-(R)-[2-(N,N-dimethylamino)ethyl]-6-[4-(4-methoxyphenylcarbonyloxy) phenylmethoxy]tetralin (330 mg) was added dropwise to THF suspension (4 ml) of lithium aluminum hydride (60 mg) under ice-cooling. 1N aqueous sodium hydroxide solution was
 25 added the reaction mixture to make the mixture basic, and the precipitate was removed by celite filtration. After the filtrate was concentrated, the residue was purified using silica gel chromatography (development solvent; ethyl acetate - methanol), and recrystallized (ethyl
 30 acetate-hexane) to give the titled compound (70 mg).

Melting point: 132 - 135°C

$[\alpha]_D^{20} = +56.9^\circ$ (c = 0.505, methanol)

Reference Example 32

2-(6-Methoxy-2-tetralinyl)-1-piperidino-1-ethanone

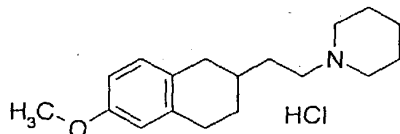


5 2-(6-Methoxy-2-tetralinyl)acetic acid (8.8 g) was dissolved in a mixed solution of THF (150 ml) and acetonitrile (50 ml), then piperidine (5.2 g), WSC (12 g), HOBT (6.0 g) and triethylamine (17 ml) were added to the solution, which was stirred at room temperature for 12
10 hours. Water was added to the reaction mixture, and extraction was conducted using ethyl acetate. The organic layer was washed with 1N hydrochloric acid, water, saturated sodium bicarbonate solution, water, and saturated aqueous sodium chloride solution, dried, and then
15 concentrated. The residue was purified using silica gel chromatography (development solvent; ethyl acetate) to give the titled compound (10.3 g). Recrystallization from hexane gave crystals of the following melting points. Melting point: 59 - 61°C

20

Reference Example 33

6-Methoxy-2-(2-piperidinoethyl)tetralin hydrochloride



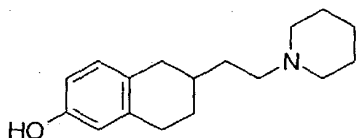
25 THF solution (50 ml) of 2-(6-methoxy-2-tetralinyl)-1-piperidino-1-ethanone (9.80 g) was added dropwise to THF suspension (100 ml) of lithium aluminum hydride (1.94 g) under ice-cooling. The temperature of the reaction mixture was raised to 60°C over 30 minutes, which was stirred for 30 minutes. After the reaction mixture was
30 cooled to room temperature, 1N aqueous sodium hydroxide solution was added to make the reaction mixture basic, and

the precipitate was removed by celite filtration. The filtrate was concentrated and the residue was made into a hydrochloride, which was then recrystallized from ethanol-IPE to give the titled compound (9.80 g).

5 Melting point: 189 - 191°C

Reference Example 34

6-Hydroxy-2-(2-piperidinoethyl)tetralin



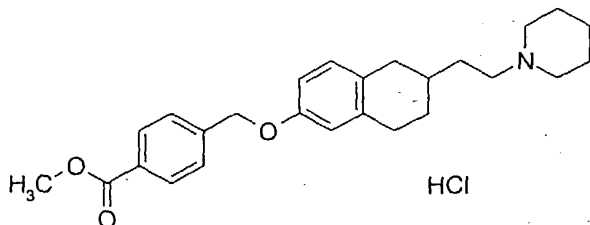
10 6-Methoxy-2-(2-piperidinoethyl)tetralin

hydrochloride (9.3 g) was added to 48% hydrobromic acid (50 ml), which was refluxed with heating for 4 hours. After the reaction mixture was concentrated under reduced pressure, saturated sodium bicarbonate solution was added to the residue to make the water layer alkaline, and the water layer was extracted using a mixed solution of THF and ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The resulting crystal was washed with IPE to give the titled compound (5.8 g).

20 Melting point: 154 - 157°C

Reference Example 35

25 Methyl 4-[[2-(2-piperidinoethyl)-6-tetralinyl]oxymethyl]benzoate hydrochloride

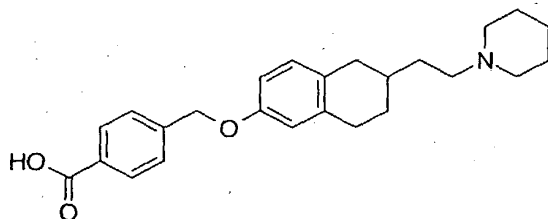


Diethyl azodicarboxylate (40% toluene solution, 5.10 g) was added dropwise to THF solution (15 ml) of 6-hydroxy-2-(2-piperidinoethyl)tetralin (1.50 g), methyl

4-(hydroxymethyl)benzoate (1.44 g), and triphenylphosphine (2.60 g) at room temperature, which was stirred for 12 hours, and then concentrated. The residue was purified using aluminum column chromatography (development solvent; hexane - hexane:ethyl acetate = 15:1), which was made into a hydrochloride. The hydrochloride was recrystallized (methanol-IPE) to give the titled compound (1.36 g).
Melting point: 190 - 193°C.

Reference Example 36

4-[[2-(2-Piperidinoethyl)-6-tetralinyl]oxymethyl]benzoic acid

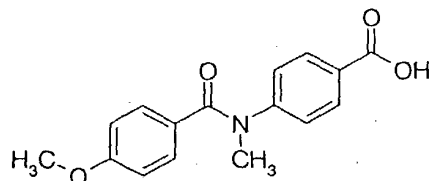


3N Aqueous sodium hydroxide solution (1.8 ml) was added to methanol solution (20 ml) of methyl 4-[[2-(2-piperidinoethyl)-6-tetralinyl]oxymethyl]benzoate hydrochloride (1.06 g), which was refluxed with heating for 6 hours. After the reaction mixture was concentrated, water was added to the reaction mixture. Further, 1N hydrochloric acid was added to make the pH of the mixture about 7. The resulting crystals were filtered to give the titled compound (0.93 g). Recrystallization from ethanol gave crystals of the following melting points.

Melting point: 105 - 108°C

Reference Example 37

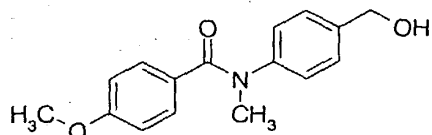
4-[N-(4-Methoxybenzoyl)-N-methylamino]benzoic acid



Aqueous solution (50 ml) of sodium carbonate (23 g) was added to THF solution (50 ml) of 4-(methylamino)benzoic acid (5.0 g), and p-anisoyl chloride (5.6 g) was added dropwise to the solution under ice-cooling, which was stirred for 15 minutes, and then stirred at room temperature for 30 minutes. Concentrated hydrochloric acid was added to the reaction mixture under ice-cooling to make the water layer acidic, and extraction was conducted using ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was purified using silica gel column chromatography (development solvent; hexane - hexane:ethyl acetate = 1:2), and recrystallized (ethyl acetate-hexane) to give the titled compound (4.8 g). Melting point: 157 - 160°C.

Reference Example 38

N-[4-(Hydroxymethyl)phenyl]-4-methoxy-N-methylbenzamide



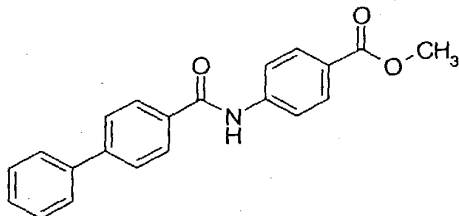
THF solution (1M, 16 ml) of borane was added dropwise to THF solution (10 ml) of 4-[N-(4-methoxybenzoyl)-N-methylamino]benzoic acid (1.14 g) under ice-cooling, which was stirred for 15 minutes, and then stirred at room temperature for 1 hour. After water was added to the reaction mixture, 1N hydrochloric acid was further added, and extraction was conducted using ethyl acetate. The organic layer was washed with water and saturated sodium bicarbonate, and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was

purified using silica gel chromatography (development solvent; hexane - hexane:ethyl acetate = 1:2), and recrystallized (ethyl acetate-hexane) to give the titled compound (770 mg).

5 Melting point: 85 - 90°C.

Reference Example 39

Methyl 4-(4-biphenylcarbonylamino)benzoate

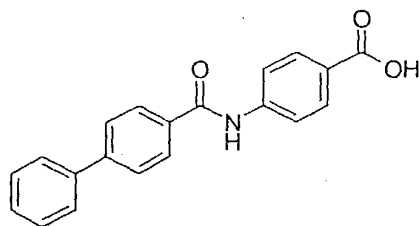


10 Oxalyl chloride (1.2 ml) and DMF (0.04 ml) were added to THF solution (30 ml) of 4-biphenylcarboxylic acid (2.184g) under ice-cooling. The reaction mixture was stirred at room temperature for 30 minutes, which was concentrated. The residue was dissolved in THF (15 ml),
15 which was added dropwise to THF solution (30 ml) of methyl 4-aminobenzoate (1.512 g) and triethylamine (2.1 ml) at 0°C. After the reaction mixture was stirred at 0°C for 30 minutes, 10% citric acid solution was added to the reaction mixture, and extraction was conducted using ethyl acetate.
20 The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The resulting crude crystal was washed with diethyl ether to give the titled compound (2.179 g).
Melting point: 247 - 251°C.

25

Reference Example 40

4-(4-Biphenylcarbonylamino)benzoic acid

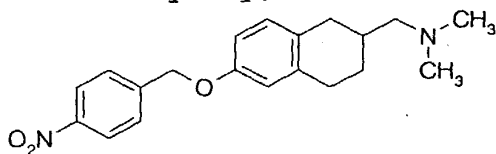


1N Aqueous sodium hydroxide solution (8 ml) was added to a mixed solution of methyl 4-(4-biphenyl)carbamoylbenzoate (1.998 g) in THF (60 ml) and methanol (20 ml), which was stirred at room temperature for 18 hours. 1N Hydrochloric acid (10 ml) was added to the reaction mixture, and extraction was conducted using ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The resulting crude crystals were washed with diethyl ether to give the titled compound (1.760 g). Melting point: >320° C.

¹H NMR (DMSO-d₆) δ: 7.37-7.57 (3H,m), 7.77 (2H,d), 7.85 (2H,d), 7.95 (4H,s), 8.08 (2H,d), 10.56 (1H,s)

Reference Example 41

2-[(N,N-Dimethylamino)methyl]-6-(4-nitrobenzyloxy)tetralin

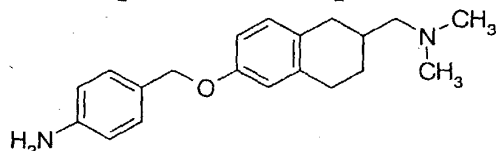


Diethyl azodicarboxylate (40% toluene solution, 9.53 g) was added dropwise to THF solution (15 ml) of 2-[(N,N-dimethylamino)methyl]-6-hydroxytetralin (1.5 g), 4-nitrobenzylalcohol (3.35 g), and triphenylphosphine (5.74 g) at room temperature, which was stirred for 24 hours. The reaction mixture was concentrated, and the residue was purified using alumina column chromatography (development solvent; hexane ~ hexane:ethyl acetate = 8:1), and recrystallized (ethyl acetate-hexane) to give the titled compound (1.29 g).

Melting point: 83 - 89°C

Reference Example 42

6-(4-Aminobenzoyloxy)-2-[(N,N-
5 dimethylamino)methyl]tetralin



After acetic acid (6 ml) was added to THF solution (12 ml) of 2-[(N,N-dimethylamino)methyl]-6-(4-nitrobenzoyloxy)tetralin (1.91 g) under ice-cooling, zinc powder (3.67 g) was further added, which was stirred for 10 6 hours. The reaction mixture was filtered, and the filtrate was concentrated. 10% aqueous potassium carbonate solution and ethyl acetate were added to the residue, the precipitate was removed by celite filtration, 15 and the filtrate was extracted using ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was purified using aluminum column chromatography (development solvent; hexane - hexane:ethyl acetate = 4:1) 20 to give the titled compound (1.05 g).

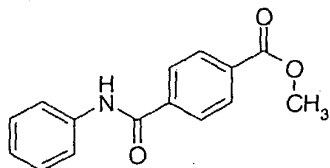
Amorphous powder:

¹H-NMR (CDCl₃) δ: 1.18-1.50(1H, m), 1.70-2.50(5H, m), 2.24(6H, s), 2.72-2.86(3H, m), 3.68(2H, brs), 4.88(2H, s), 6.58-6.82(4H, m), 6.99(1H, s), 7.14-7.30(2H, m).

25

Reference Example 43

Methyl 4-anilinocarbonylbenzoate



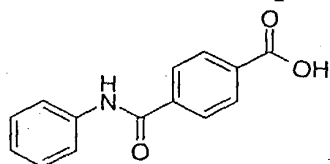
4-Methoxycarbonyl benzoic acid (540 mg), aniline 30 (0.27 ml), WSC (863 mg) and triethylamine (0.84 ml) were

added to THF (20 ml). After the reaction mixture was stirred at room temperature for 20 hours, the reaction mixture was placed in water, and extraction was conducted using ethyl acetate-THF (1:1). The organic layer was washed with water, saturated sodium bicarbonate solution, and saturated aqueous sodium chloride solution, dried, and then concentrated. The resulting crude crystals were recrystallized from ethyl acetate-hexane to give the titled compound (659 mg).

10 Melting point: 189 - 190°C

Reference Example 44

4-Anilinocarbonylbenzoic acid



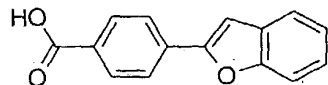
15 8 mol of aqueous sodium hydroxide solution (8 ml) was added to methanol (16 ml) - THF (6 ml) solution of 4-methyl anilinocarbonylbenzoate (511 mg), which was stirred at room temperature for 1 hour. 1 mol of hydrochloric acid was added to the reaction mixture to make the pH of the mixture to 5, extraction was conducted using ethyl acetate-THF (1:1). The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The resulting residue was washed with hexane to give the titled compound (480 mg).

20

25 Melting point: 305 - 307°C.

Reference Example 45

4-(2-Benzo[b]furanyl)benzoic acid



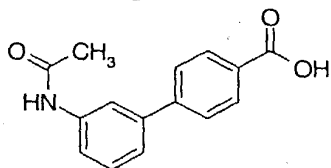
30 Benzo[*b*]furanyl-2-boric acid (2.1 g), palladium tetratriphenylphosphine (200 mg) and 2M aqueous sodium

carbonate solution were added to toluene (40 ml) - ethanol (10 ml) solution of ethyl 4-bromobenzoate (2.3 g), which was refluxed at 80°C for 5 hours under an argon atmosphere.

The reaction mixture was diluted with water, and
5 extraction was conducted using ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The resulting residue was purified using silica gel
10 chromatography (development solvent; ethyl acetate:hexane = 1:4), and concentrated, which was dissolved in methanol (10 ml) - THF (10 ml). 8 mol of aqueous sodium hydroxide solution (8 ml) was added to the resulting solution at room temperature, which was stirred for 2 hours. After 1 mol of hydrochloric acid was added to the reaction mixture to
15 make the mixture acidic, extraction was conducted using ethyl acetate-THF (1:1). The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The resulting residue was washed with hexane to give the titled compound (2.272 g).
20 Melting point: 292 - 294°C.

Reference Example 46

3'-Acetylamino-4-biphenylylcarboxylic acid

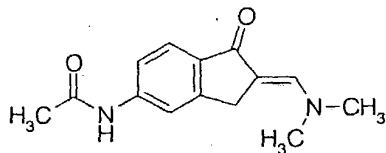


25 The titled compound was produced in the same manner as in Reference Example 45.

Melting point: 300 - 301°C

Reference Example 47

30 N-[2-[(E)-(Dimethylamino)methylidene]-1-oxo-2,3-dihydro-1H-inden-5-yl]acetamide

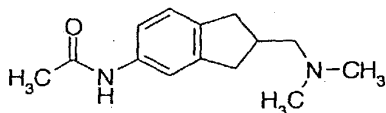


Dimethylformamide dimethylacetal was added to 5-acetamido-1-indanone (2.5 g, 13.2 mmol), which was stirred at 100°C for 3.5 hours, and cooled to room temperature. The precipitated crude products were collected, which was washed with ethyl acetate to give the titled compound (2.73 g).

¹H NMR (DMSO-d₆) δ: 2.08 (3H, s), 3.13 (6H, s), 3.87 (2H, s), 7.31 (1H, s), 7.52 (2H, m), 7.86 (1H, s), 10.16 (1H, s).

Reference Example 48

N-[2-[(Dimethylamino)methyl]-2,3-dihydro-1H-inden-5-yl]acetamide



N-[2-[(E)-(Dimethylamino)methylidene]-1-oxo-2,3-dihydro-1H-inden-5-yl]acetamide (2.70 g, 12.3 mmol) obtained in Reference Example 47 and 10% palladium-carbon (0.3 g) were added to a mixed solution of methanol (60 ml) and acetic acid (6 ml), which was stirred at 40°C under a hydrogen atmosphere for 1 day. After the catalyst was filtered, the filtrate was distilled out under reduced pressure. 1N hydrochloric acid (15 ml) was added to the reaction mixture, which was washed with ethyl acetate.

Then, potassium carbonate was added to the mixture, and extraction was conducted using ethyl acetate. The extract was washed with saturated aqueous sodium chloride solution, dried using anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. The resulting residue was purified using aluminum column chromatography (development solvent: ethyl acetate) to give the titled

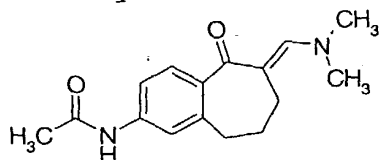
compound.

^1H NMR (CDCl_3) δ : 2.15 (3H, s), 2.25 (6H, s), 2.28 (2H, m), 2.61 (3H, m), 3.02 (2H, m), 7.11 (2H, m), 7.26 (1H, s), 7.39 (1H, s).

5

Reference Example 49

N-[6-[(E)-(Dimethylamino)methylidene]-5-oxo-6,7,8,9-tetrahydro-5H-benzo[a]cyclohepten-2-yl]acetamide



10 The titled compound was obtained by carrying out the same operation as in Reference Example 47, using N-(5-oxo-6,7,8,9-tetrahydro-5H-benzo[a]cyclohepten-2-yl)acetamide.

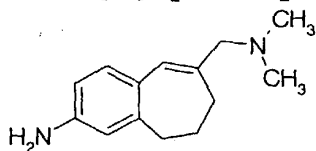
^1H -NMR (CDCl_3) δ : 1.78-1.90 (2H, m), 2.17 (3H, s), 2.34 (2H, t, J = 6.6 Hz), 2.74 (2H, t, J = 6.8 Hz), 3.11 (6H, s), 7.21 (1H, d, J = 8.1 Hz), 7.48-7.63 (3H, m), 7.73 (1H, s).

15

Melting point: 177 - 180°C (crystallization solvent: ethyl acetate-diethyl ether)

20 Reference Example 50

8-[(Dimethylamino)methyl]-6,7-dihydro-5H-benzo[a]cyclohepten-3-amine



25 The titled compound was obtained as an oily substance by carrying out the same operation as in Example 41-2), using N-[6-[(E)-(dimethylamino)methylidene]-5-oxo-6,7,8,9-tetrahydro-5H-benzo[a]cyclohepten-2-yl]acetamide obtained in Reference Example 49.

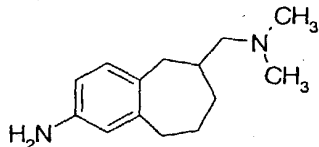
^1H -NMR (CDCl_3) δ : 1.90-2.01 (2H, m), 2.22 (6H, s), 2.35 (2H, t, J = 6.3 Hz), 2.72 (2H, t, J = 5.4 Hz), 2.91 (2H, s), 3.7

30

(2H, br, NH₂), 6.28 (1H, s), 6.40-6.50 (2H, m), 6.94 (1H, d, J = 7.8 Hz).

Reference Example 51

- 5 6-[(Dimethylamino)methyl]-6,7,8,9-tetrahydro-5H-benzo[a]cyclohepten-2-amine

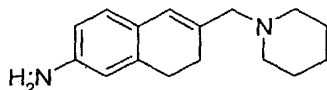


- The titled compound was obtained as an oily substance, by carrying out the same operation as in Reference Example 10 48, using 8-[(dimethylamino)methyl]-6,7-dihydro-5H-benzo[a]cyclohepten-3-amine.
- ¹H-NMR (CDCl₃) δ: 1.30-1.63 (3H, m), 1.65-2.22 (10H, m), 2.44-2.80 (4H, m), 3.5 (2H, br, NH₂), 6.35-6.48 (2H, m), 6.92 (1H, d, J = 7.8 Hz).

15

Reference Example 52

- 6-(1-Piperidinylmethyl)-7,8-dihydro-2-naphthalenamine



- 1) A mixture of 6-acetamido-2-(N,N-dimethylaminomethylidene)-1-tetralone (11 g) obtained in 20 Example 41-1) and piperidine (100 ml) was refluxed with heating for 24 hours. After excess piperidine was distilled out under reduced pressure, the resulting residue was crystallized using tetrahydrofuran-isopropyl ether to 25 give 6-acetamido-2-(1-piperidinylmethylidene)-1-tetralone (7 g) as a light yellow powder.

- 2) The titled compound was obtained as an amorphous powder by carrying out the same operations as in Example 41-2), using 6-acetamido-2-(1-piperidinylmethylidene)-1-tetralone obtained in above 1).

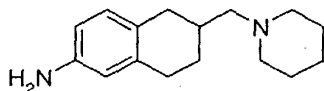
30

¹H-NMR (CDCl₃) δ: 1.44-1.57 (6H, m), 2.25-2.34 (6H, m), 2.72 (2H, t, J=8.0 Hz), 2.98 (2H, s), 3.59 (2H, s), 6.23 (1H,

s), 6.45-6.47 (2H, m), 6.81 (1H, d, J=8.7 Hz).

Reference Example 53

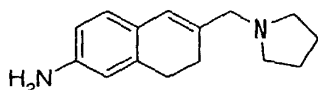
5 6-(1-Piperidinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenamine



The titled compound was obtained as an amorphous powder by carrying out the same operations as in Reference Example 48, using 6-(1-piperidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 52.
10 ¹H NMR (CDCl₃) δ: 1.25-2.82 (19H, m), 3.36 (2H, bs), 6.44-6.49 (2H, m), 6.88 (1H, d, J=8.1 Hz).

Reference Example 54

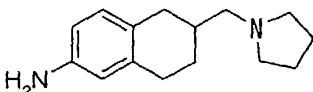
15 6-(1-Pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine



The titled compound was obtained as an amorphous powder by carrying out the same operations as in Reference Example 52, using 6-acetamido-2-(N,N-dimethylaminomethylidene)-1-tetralone obtained in
20 Example 41-1).
¹H NMR (CDCl₃) δ: 1.76-1.80 (4H, m), 2.30 (2H, t, J = 7.8 Hz), 2.47-2.49 (4H, m), 2.74 (2H, t, J = 7.8 Hz), 3.13 (2H, s), 3.59 (2H, brs), 6.26 (1H, s), 6.45-6.47 (2H, m),
25 6.82 (1H, d, J = 8.6Hz).

Reference Example 55

6-(1-Pyrrolidinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenamine



30

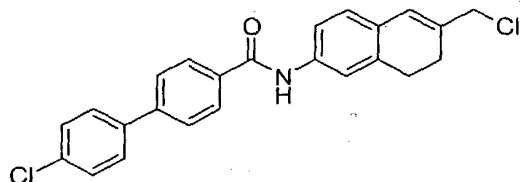
The titled compound was obtained as an amorphous

powder by carrying out the same operations as in Reference Example 48, using 6-(1-pyrrolidinymethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

¹H NMR (CDCl₃) δ : 1.45-1.90 (1H, m), 1.55-2.80 (16H, m),
 5 3.48 (2H, brs), 6.44 (1H, s), 6.47 (2H, d, J = 8.1 Hz),
 6.88 (2H, d, J = 8.1 Hz).

Reference Example 56

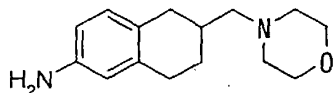
4'-Chloro-N-[6-(chloromethyl)-7,8-dihydro-2-naphthalenyl] [1,1'-biphenyl]-4-carboxamide



After 1-chloroethyl chloroformate (0.23 ml) was added to tetrahydrofuran solution (30 ml) of 4'-chloro-N-[6-(dimethylamino)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide (750 mg) at
 15 -78°C, the temperature of the solution was raised to room temperature over 30 minutes. The solvent was distilled out under reduced pressure. The resulting residue was crystallized using tetrahydrofuran-n-hexane to give the
 20 titled compound (600 mg).
 Melting point: 179 - 181°C (crystallization solvent: tetrahydrofuran-n-hexane)

Reference Example 57

25 6-(4-Morpholinymethyl)-5,6,7,8-tetrahydro-2-naphthalenamine



The titled compound was obtained as an amorphous powder by carrying out, in order, the same operations as
 30 in Reference Example 52 and Reference Example 48, using 6-acetamido-2-(N,N-dimethylaminomethylidene)-1-

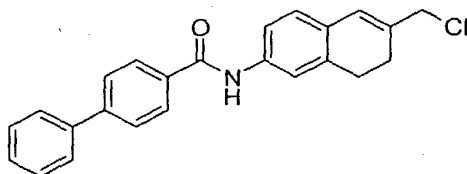
tetralone obtained in Example 41-1).

^1H NMR (CDCl_3) δ : 1.22-1.41 (1H, m), 1.80-1.82 (2H, m), 2.22-2.34 (10H, m), 3.50 (2H, s), 3.69-3.72 (1H, m), 6.40 (1H, s), 6.44 (1H, d, $J = 8.1$ Hz), 6.85 (1H, d, $J = 8.1$ Hz).

5

Reference Example 58

N-[6-(Chloromethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



10

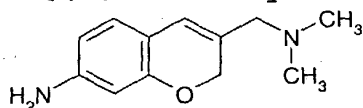
The titled compound was obtained by carrying out the same operations as in Reference Example 56, using N-[6-((dimethylamino)methyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide obtained in Example 47.

15

Melting point: 163 - 165°C (crystallization solvent: tetrahydrofuran-n-hexane)

Reference Example 59

3-[(N,N-Dimethylamino)methyl]-2H-chromen-7-amine



20

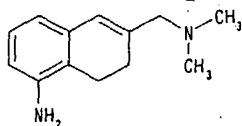
The titled compound was obtained by carrying out, in order, the same operations as in Examples 41-1) and 41-2), using 7-acetylamino-3,4-dihydrochromen-4-on.

^1H -NMR (CDCl_3) δ : 2.20 (6H, s), 2.94 (2H, s), 3.66 (2H, brs), 4.71 (2H, s), 6.16-6.21 (2H, m), 6.76 (1H, d, $J = 7.8$ Hz).

25

Reference Example 60

6-[(Dimethylamino)methyl]-7,8-dihydro-1-naphthalenamine



1) Methyl 4-(2-aminophenyl)butanoate hydrochloride (7.20 g, 0.037 mol) synthesized by a known method by documents (Synthetic communications, 26(18), 3443 (1996)) and triethylamine (5.06 g, 0.05 mol) were dissolved in tetrahydrofuran (60ml). Acetyl chloride (3.51 g, 0.045 mol) was added dropwise to the mixture, which was stirred at room temperature for 30 minutes. Ethyl acetate and 1N hydrochloric acid were added to the reaction mixture, and extraction was conducted. The organic layer was washed with water, concentrated and dried. A mixed solution of ethyl acetate - n-hexane (1 : 1) was added to the residue. The crystallized product was collected by filtration, to give methyl 4-(2-acetylamino-phenyl)butanoate (6.40g) as a white powder.

¹H-NMR (CDCl₃) δ: 1.77-1.86 (2H, m), 2.29 (3H, s), 2.41-2.45 (2H, m), 2.59-2.62 (2H, m), 3.74 (3H, s), 7.03 (1H, t, J=7.3 Hz), 7.11-7.12 (1H, m), 7.22 (1H, t, J=7.3 Hz), 8.08 (1H, d, J=8.1 Hz), 8.33 (1H, s).

2) Polyphosphoric acid (100g) was heated at 130°C, then methyl 4-(2-acetylamino-phenyl)butanoate (6.40g, 0.027mol) obtained in 1) was added under stirring. After stirring for 1 hour, the reaction mixture was poured into ice water, and ethyl acetate and water were added, then extraction was conducted by adding water. The organic layer was washed with saturated sodium hydrogen carbonate solution and aqueous sodium chloride solution, and concentrated. A mixed solution of ethyl acetate - n-hexane (1:1) was added to the residue, and the crystallized product was collected by filtration, to give 5-acetylamino-1-tetralone (2.80g) as a white powder.

¹H-NMR (CDCl₃) δ: 2.10-2.19 (2H, m), 2.24 (3H, s), 2.66 (2H, t, J=6.3 Hz), 2.84 (2H, t, J=5.7 Hz), 7.06 (1H, brs), 7.34 (1H, t, J=7.5 Hz), 7.82 (1H, d, J=7.5 Hz), 7.95 (1H, d, J=7.5 Hz).

3) 5-Acetylamino-1-tetralone (0.6g, 3.0 mmol) obtained was dissolved in dimethylformamide dimethylacetal

(20ml), which was refluxed with heating for 4 hours. The crystallized product was collected by filtration, which was washed with ethyl acetate, to give 5-acetylamino-2-(dimethylamino)methylidene-1-tetralone (0.58g) as a yellow powder.

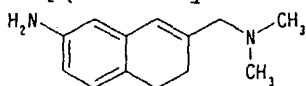
$^1\text{H-NMR}$ (CDCl_3) δ : 2.21 (3H, s), 2.68-2.72 (2H, m), 2.86-2.90 (2H, m), 3.11 (6H, s), 7.26-7.31 (2H, m), 7.62 (1H, m), 7.69 (1H, s), 7.92 (1H, m).

4) Sodium triacetoxyhydroborate (424 mg, 2.0 mmol) was dissolved in a mixed solution of ethyl acetate (5ml) and tetrahydrofuran (1ml) under ice-cooling. 5-Acetylamino-2-dimethylaminomethylidene-1-tetralone (129 mg, 0.5 mmol) obtained in 3) was added to the mixture, which was stirred for 15 minutes. The reaction mixture was concentrated, and methanol (10ml) was added to the residue, and sodium borohydride (38 mg, 1 mmol) was added under ice-cooling. After stirring for 1 hour, the reaction mixture was concentrated. 5N Hydrochloric acid and ethyl acetate were added to the residue, and extraction was conducted. The water layer was refluxed with heating for 2 hours. 4N sodium hydroxide solution and ethyl acetate were added to the reaction mixture, and extraction was conducted. The organic layer was washed with water, and concentrated. The residue was purified by alumina column chromatography (development solvent; ethyl acetate : n-hexane=1:1), to give the titled compound (80 mg) as a colorless oily substance.

$^1\text{H-NMR}$ (CDCl_3) δ : 2.24(6H, s), 2.37(2H, t, $J=8.1$ Hz), 2.63(2H, t, $J=8.1$ Hz), 2.97(2H, s), 3.58(2H, brs), 6.29(1H, s), 6.53(1H, d, $J=8.1$ Hz), 6.57 (1H, d, $J=8.1$ Hz), 6.97(1H, t, $J=8.1$ Hz).

Reference Example 61

7-[(Dimethylamino)methyl]-5,6-dihydro-2-naphthalenamine



1) 7-Nitro-1-tetralone (8.32 g, 0.044 mol) and concentrated hydrochloric acid (24 ml, 0.29 mol) were dissolved in methanol (100 ml), and an iron powder (7.30 g, 0.13 mol) was gradually added over 1 hour. After stirring for 1 hour, the reaction mixture was concentrated.

4N Sodium hydroxide solution and ethyl acetate were added to the residue, and extraction was conducted. The organic layer was dried, and concentrated. Tetrahydrofuran (100 ml) and triethylamine (5.05 g, 0.05 mol) was added to the residue. Further, acetyl chloride (3.92 g, 0.05 mol) was added under ice-cooling. After stirring for 30 minutes, ethyl acetate and 1N hydrochloric acid were added, and extraction was conducted. The organic layer was concentrated, and the residue was purified with silica gel column chromatography (development solvent: ethyl acetate), to give 7-acetylamino-1-tetralone (7.52 g) as a white powder.

$^1\text{H-NMR}$ (CDCl_3) δ : 2.09-2.18 (2H, m), 2.21 (3H, s), 2.65 (2H, t, $J=6.3$ Hz), 2.94 (2H, t, $J=6.3$ Hz), 7.24 (1H, d, $J=8.4$ Hz), 7.82 (1H, s), 7.98 (1H, brs), 8.15 (1H, d, $J=7.5$ Hz).

2) 7-Acetylamino-2-[(dimethylamino)methylidene]-1-tetralone (2.95 g) was obtained as a white powder by the same method as in Reference Example 60-3), using 7-acetylamino-1-tetralone (3.00 g, 0.0148 mol) obtained in 1).

$^1\text{H-NMR}$ (CDCl_3) δ : 2.17 (3H, s), 2.78-2.82 (2H, m), 2.88-2.93 (2H, m), 3.14 (6H, s), 7.14 (1H, d, $J=8.1$ Hz), 7.74 (1H, s), 7.76 (1H, s), 8.09-8.12 (1H, m), 8.24 (1H, s).

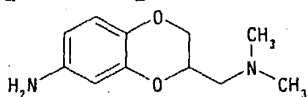
3) The titled compound (300 mg) was obtained as a colorless oily substance by the same method as in Reference Example 60-4), using 7-acetylamino-2-[(dimethylamino)methylidene]-1-tetralone (628 mg, 2.43 mmol) obtained in 2).

$^1\text{H-NMR}$ (CDCl_3) δ : 2.23 (6H, s), 2.29 (2H, t, $J=8.4$ Hz), 2.71 (2H, t, $J=8.4$ Hz), 2.97 (2H, s), 3.52 (2H, brs), 6.24 (1H, s), 6.41 (1H, s), 6.46 (1H, d, $J=7.8$ Hz), 6.90 (1H, d,

J=7.8 Hz).

Reference Example 62

5 N,N-Dimethyl-N-[(7-amino-2,3-dihydro-1,4-benzodioxin-2-yl)methyl]amine



1) 1,2-Dihydroxy-4-nitrobenzene (5.00 g, 0.032 mol), potassium carbonate (9.67 g, 0.07 mol) and epibromohydrin (5.30 g, 0.039 mol) were dissolved in dimethylformamide (100ml), which was stirred at 100°C for 1 hour. Water was added to the reaction mixture, and extraction was conducted using ethyl acetate. The organic layer was washed with water, and concentrated. The residue was purified by alumina column chromatography (development solvent : ethyl acetate). The eluent was washed with a mixed solution of ethyl acetate - n-hexane (1:1), to give (7-nitro-2,3-dihydro-1,4-benzodioxin-2-yl)methanol (3.31 g) as a white powder.

¹H-NMR (CDCl₃) δ : 1.95-1.99 (1H, m), 3.89-3.97 (2H, m), 4.19-4.29 (2H, m), 4.41-4.45 (1H, m), 6.96 (1H, d, J=8.6 Hz), 7.78-7.81 (2H, m).

2) (7-Nitro-2,3-dihydro-1,4-benzodioxin-2-yl)methanol (1.00 g, 4.74 mmol) obtained in 1) and triethylamine (719 mg, 7.10 mmol) were dissolved in dimethylformamide (30 ml), and methanesulfonyl chloride (651 mg, 5.68 mmol) was added, which was stirred at room temperature for 30 minutes. Then, an aqueous dimethylamine solution was added and stirred at 60°C for 5 hours. Water was added to the reaction mixture, and extraction was conducted using ethyl acetate. The organic layer was washed with water, and concentrated. The residue was purified by alumina column chromatography (development solvent ; ethyl acetate : n-hexane = 3:7), to give N,N-dimethyl-N-[(7-nitro-2,3-dihydro-1,4-benzodioxin-2-yl)methyl]amine (802 mg) as a colorless oily substance.

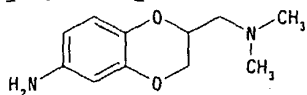
$^1\text{H-NMR}$ (CDCl_3) δ : 2.34 (6H, s), 2.50-2.68 (2H, m), 4.02-4.09 (2H, m), 4.30-4.36 (1H, m), 4.39-4.44 (2H, m), 6.94 (1H, d, $J=8.9\text{Hz}$), 7.76-7.84 (2H, m).

3) N,N-Dimethyl-N-[(7-nitro-2,3-dihydro-1,4-benzodioxin-2-yl)methyl]amine (802 mg, 3.37 mmol) obtained in 2) and concentrated hydrochloric acid (3 ml) was dissolved in methanol (10 ml), and an iron powder (0.80 g, 14 mmol) was quietly added over 1 hour. After stirring for 1 hour, the reaction mixture was concentrated. 4N Sodium hydroxide solution and ethyl acetate were added to the residue, and extraction was conducted. The organic layer was dried, and concentrated. The residue was purified by silica gel column chromatography (development solvent: ethyl acetate - n-hexane = 3:7), to give the titled compound (514 mg) as a colorless oily substance.

$^1\text{H-NMR}$ (CDCl_3) δ : 2.32 (6H, s), 2.43-2.64 (2H, m), 3.40 (2H, s), 3.86-3.93 (1H, m), 4.19-4.27 (2H, m), 6.18-6.22 (1H, m), 6.29 (1H, s), 6.67 (1H, d, $J=8.7\text{ Hz}$).

Reference Example 63

N,N-Dimethyl-N-[(6-amino-2,3-dihydro-1,4-benzodioxin-2-yl)methyl]amine



1) 1,2-Dihydroxy-4-nitrobenzene (4.65 g, 0.030 mol), potassium carbonate (8.71 g, 0.063 mol) and methoxymethyl chloride (2.42 g, 0.030 mol) were dissolved in dimethylformamide (50 ml), which was stirred at 40°C for 30 minutes. Epibromohydrin (7.20 g, 0.045 mol) was added to the mixture, which was stirred at 60°C for 80 minutes. Then water was added, and extraction was conducted using ethyl acetate. The organic layer was washed with water, and concentrated. The residue was purified by alumina column chromatography (development solvent: ethyl acetate - n-hexane = 1:4), to give 2-[[2-(methoxymethoxy)-5-nitrophenoxy]methyl]oxirane (2.61 g) as a white powder.

$^1\text{H-NMR}$ (CDCl_3) δ : 2.79-2.81 (1H, m), 2.93-2.96 (1H, m), 3.41 (1H, m), 3.53 (3H, s), 4.01-4.07 (1H, m), 4.40-4.45 (1H, m), 5.32 (2H, s), 7.22 (1H, d, $J=9.0$ Hz), 7.82-7.91 (2H, m).

5 2) 2-[[2-(Methoxymethoxy)-5-nitrophenoxy]methyl]oxirane (4.00 g, 0.016 mol) obtained in 1) was dissolved in methanol (50 ml), and 10% hydrochloric acid-methanol solution (10 ml) was added, which was stirred at room temperature for 30 minutes. The
10 solvent was concentrated, and methanol (30 ml) and potassium carbonate (6.50 g, 0.047 mol) were added to the residue, which was stirred at 60°C for 1 hour. The solvent was concentrated, water was added, and extraction was conducted using ethyl acetate. The organic layer was
15 washed with water, and concentrated. The residue was purified by alumina column chromatography (development solvent; ethyl acetate), to give (6-nitro-2,3-dihydro-1,4-benzodioxin-2-yl)methanol (2.12 g) as a white powder. $^1\text{H-NMR}$ (CDCl_3) δ : 1.90-1.94 (1H, m), 3.89-3.97 (2H, m),
20 4.19-4.28 (2H, m), 4.41-4.45 (1H, m), 6.97 (1H, d, $J=8.6$ Hz), 7.78-7.82 (2H, m).

3) N,N-Dimethyl-N-[(6-nitro-2,3-dihydro-1,4-benzodioxin-2-yl)methyl]amine (910 mg) was obtained as a colorless oily substance, by the same method as in Reference
25 Example 62-2), using (6-nitro-2,3-dihydro-1,4-benzodioxin-2-yl)methanol (1.00 g, 4.74 mmol) obtained in 2).

$^1\text{H-NMR}$ (CDCl_3) δ : 2.35 (6H, s), 2.52-2.70 (2H, m), 3.98-4.05 (2H, m), 4.35-4.39 (3H, m), 6.95-6.98 (1H, m), 7.77-7.80
30 (2H, m).

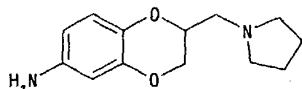
4) The titled compound (750 mg) was obtained as a colorless oily substance, by the same method as in Reference Example 62-3), using N,N-dimethyl-N-[(6-nitro-2,3-dihydro-1,4-benzodioxin-2-yl)methyl]amine (910 mg, 3.82
35 mmol) obtained in 3).

$^1\text{H-NMR}$ (CDCl_3) δ : 2.32 (6H, s), 2.43-2.64 (2H, m), 3.40 (2H,

s), 3.86-3.92 (1H, m), 4.13-4.27 (2H, m), 6.19-6.28 (2H, m), 6.67-6.70 (1H, m).

Reference Example 64

- 5 1-[(6-Amino-2,3-dihydro-1,4-benzodioxin-2-yl)methyl]pyrrolidine

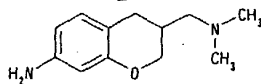


- 1) 1-[(6-Nitro-2,3-dihydro-1,4-benzodioxin-2-yl)methyl]pyrrolidine (1.30 g) was obtained as a colorless oily substance, by the same method as in Reference Example 62-2), using (6-nitro-2,3-dihydro-1,4-benzodioxin-2-yl)methanol (1.12 g, 5.30 mmol) and pyrrolidine (10 ml).
 10 ¹H-NMR (CDCl₃) δ: 1.79-1.83 (4H, m), 2.60-2.62 (4H, m), 2.78 (2H, d, J=5.9 Hz), 4.00-4.07 (1H, m), 4.38-4.42 (2H, m),
 15 6.95-6.98 (1H, m), 7.76-7.80 (2H, m).

- 2) The titled compound (1.03 g) was obtained as a colorless oily substance, by the same method as in Reference Example 62-3), using 1-[(6-nitro-2,3-dihydro-1,4-benzodioxin-2-yl)methyl]pyrrolidine (1.30 g, 4.92 mmol).
 20 ¹H-NMR (CDCl₃) δ: 1.74-1.83 (4H, m), 2.54-2.63 (4H, m), 2.69-2.72 (2H, m), 3.40 (2H, s), 3.91-3.97 (1H, m), 4.18-4.30 (2H, m), 6.18-6.25 (2H, m), 6.70 (1H, d, J=8.4 Hz).

- 25 Reference Example 65

N-[(7-Amino-3,4-dihydro-2H-chromen-3-yl)methyl]-N,N-dimethylamine



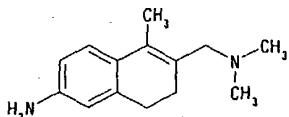
- 3-[(N,N-Dimethylamino)methyl]-2H-chromen-7-amine
 30 (150 mg, 0.73 mmol) obtained in Reference Example 59, 1N hydrochloric acid (0.5 ml) and 10% palladium carbon (40 mg) was dissolved in methanol (5 ml), and catalytic hydrogenation was conducted under normal temperature and normal pressure. After a catalyst was filtered out, the

filtrate was concentrated, and the residue was purified by alumina column chromatography (development solvent; ethyl acetate : n-hexane = 3:7), to give the titled compound (15 mg) as a colorless oily substance.

- 5 $^1\text{H-NMR}$ (CDCl_3) δ : 2.20-2.24 (3H, m), 2.24(6H, m), 2.30-2.40 (1H, m), 2.75-2.80 (1H, m), 3.60 (1H, m), 3.75-3.80 (2H, m), 4.20-4.25 (1H, m), 6.20 (1H, m), 6.21-6.25 (1H, m), 6.82 (1H, d, $J=7.8$ Hz).

10 Reference Example 66

6-[(Dimethylamino)methyl]-5-methyl-7,8-dihydro-2-naphthalenamine



- 15 1) 6-Acetylamino-1-tetralone (5.5 g, 0.027 mol) and dimethylmethylenammonium chloride (6.3 g, 0.068 mol) were dissolved in a mixed solution of acetonitrile (100 ml) and tetrahydrofuran (100 ml), which was stirred for 48 hours.

The crystallized product was collected by filtration, washed with tetrahydrofuran, and dissolved in ethyl acetate. 0.5N Sodium hydroxide solution was added to the solution for liquid separation. The organic layer was concentrated, to give 6-acetylamino-2-
20 [(dimethylamino)methyl]-1-tetralone (4.48 g) as a colorless oily substance.

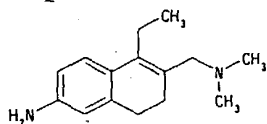
- 25 2) 6-Acetylamino-2-[(dimethylamino)methyl]-1-tetralone (260 mg, 1.00 mmol) obtained was dissolved in tetrahydrofuran (10 ml). 1M Methyl magnesium bromide - tetrahydrofuran solution (3 ml)(3.00 mmol) was added to the solution under ice-cooling, which was stirred at room
30 temperature for 16 hours. Aqueous ammonium chloride solution was added to the reaction mixture, and extraction was conducted using ethyl acetate. The organic layer was concentrated, and 5N hydrochloric acid and ethyl acetate were added to the residue for liquid separation.

Concentrated hydrochloric acid was added to the water layer, which was refluxed for 4 hours. The reaction mixture was concentrated, and 1N sodium hydroxide solution and ethyl acetate were added to the residue and extraction was conducted. The organic layer was concentrated, and the residue was purified by alumina column chromatography (development solvent; ethyl acetate : n-hexane = 3:7), to give the titled compound (83 mg) as a colorless oily substance.

¹H-NMR (CDCl₃) δ: 2.04 (3H, s), 2.24 (6H, s), 2.28 (2H, t, J=7.4 Hz), 2.66 (2H, t, J=7.4 Hz), 3.04 (2H, s), 3.62 (2H, s), 6.49 (1H, s), 6.51-6.55 (1H, m), 7.10 (1H, d, J=8.1 Hz).

Reference Example 67

6-[(Dimethylamino)methyl]-5-ethyl-7,8-dihydro-2-naphthalenamine

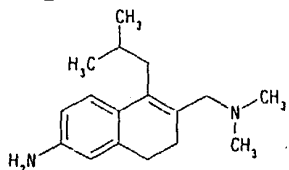


The titled compound was obtained as a colorless oily substance by the same manner as in Reference Example 66-2), using 6-acetylamino-2-(dimethylamino)methyl-1-tetralone obtained in Reference Example 66-1) and ethyl magnesium bromide.

¹H-NMR (CDCl₃) δ: 1.06 (3H, t, J=7.5 Hz), 2.24 (6H, s), 2.27 (2H, m), 2.52-2.66 (4H, m), 3.04 (2H, s), 3.61 (2H, s), 6.51 (1H, s), 6.51-6.55 (1H, m), 7.11 (1H, d, J=8.1 Hz).

Reference Example 68

6-[(Dimethylamino)methyl]-5-isobutyl-7,8-dihydro-2-naphthalenamine



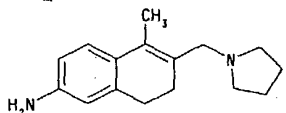
The titled compound was obtained as a colorless oily substance by the same manner as in Reference Example 66-2), using 6-acetylamino-2-[(dimethylamino)methyl]-1-tetralone obtained in Reference Example 66-1) and isobutyl magnesium bromide.

¹H-NMR (CDCl₃) δ: 0.88 (6H, d, J=6.7 Hz), 1.73-1.79 (1H, m), 2.21 (6H, s), 2.28 (2H, t, J=7.0 Hz), 2.44 (2H, d, J=7.3 Hz), 2.63 (2H, t, J=7.0 Hz), 3.09 (2H, s), 3.60 (2H, s), 6.49 (1H, s), 6.51-6.53 (1H, m), 7.08 (1H, d, J=7.8 Hz).

10

Reference Example 69

5-Methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine



1) 6-Acetylamino-2-[(dimethylamino)methylidene]-1-tetralone (4.90 g, 0.017 mol) obtained in Example 41-1) was suspended in pyrrolidine (25 ml), which was refluxed with heating for 2 hours. The crystallized product was collected by filtration, washed with a mixed solution of ethyl acetate and n-hexane (1:1), to give 6-acetylamino-2-(1-pyrrolidinylmethylidene)-1-tetralone (5.03 g) as yellow crystals.

¹H-NMR (CDCl₃) δ: 1.75-2.00 (4H, m), 2.19 (3H, s), 2.70-3.00 (4H, m), 3.50-3.70 (4H, m), 7.20-7.25 (1H, m), 7.67 (1H, s), 7.70-7.90 (2H, m), 7.97 (1H, d, J=8.4 Hz).

2) Sodium triacetoxyhydroborate (3.18 g, 0.015 mol) was dissolved in a mixed solution of ethyl acetate (50 ml) and tetrahydrofuran (12.5 ml) under ice-cooling, and 6-acetylamino-2-(1-pyrrolidinylmethylidene)-1-tetralone (2.84 g, 0.01 mol) obtained in 1) was added. After stirring for 1 hour, the reaction mixture was concentrated. 1N Sodium hydroxide solution and ethyl acetate were added to the residue, which was stirred. The crystallized product was collected by filtration, washed with a mixed solution

of ethyl acetate and n-hexane (1:1), to give 6-acetylamino-2-(1-pyrrolidinylmethyl)-1-tetralone (2.65 g) as a white powder.

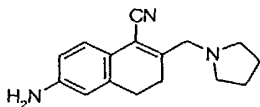
¹H-NMR (CDCl₃) δ: 1.78 (4H, m), 1.90-2.02 (1H, m), 2.20 (3H, s), 2.35-2.98 (10H, m), 7.20-7.23 (1H, m), 7.57 (1H, s), 7.66 (1H, m), 7.97 (1H, d, J=8.4 Hz).

3) The titled compound was obtained by the same manner as in Reference Example 66-2), using 6-acetylamino-2-(1-pyrrolidinylmethyl)-1-tetralone obtained in 2).

¹H-NMR (CDCl₃) δ: 1.73-1.79 (4H, m), 2.04 (3H, s), 2.31 (2H, t, J=7.4 Hz), 2.49-2.54 (4H, m), 2.65 (2H, t, J=7.8 Hz), 3.24 (2H, s), 3.60 (2H, brs), 6.48-6.54 (2H, m), 7.09 (1H, d, J=8.1 Hz).

Reference Example 70

6-Amino-2-(1-pyrrolidinylmethyl)-3,4-dihydro-1-naphthalenecarbonitrile



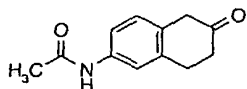
Trimethylsilylnitrile (1.02 ml, 7.68 mmol) and zinc iodide (22 mg, 0.0698 mmol) were added to dichloroethane solution (9 ml) of 6-acetylamino-2-(1-pyrrolidinylmethyl)-1-tetralone (1.00 g, 3.49 mmol) obtained in Reference Example 69-2), which was stirred at room temperature for 2 days. The solvent was distilled out under reduced pressure. Ethyl acetate was added to the obtained oily substance, which was washed with saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. The resulting residue was purified by alumina column chromatography (development solvent; ethyl acetate), to give trimethylsilylcyanohydrin form (1.21 g) as an oily substance. 2.5N Hydrochloric acid was added to the oily substance (978 mg, 2.73 mmol), which was stirred at 100°C for 1.5 hours. The aqueous solution obtained was

washed with ethyl acetate. Potassium carbonate was added to the water layer to make it alkaline, and extraction was conducted using ethyl acetate. The extract was washed with saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and the solvent was distilled out under reduced pressure. The resulting oily substance was purified by alumina column chromatography (development solvent; hexane:ethyl acetate = 5:1), to give the titled compound (358 mg).

¹H NMR (CDCl₃) δ: 1.80 (4H, m), 2.56 (6H, m), 3.73 (2H, m), 3.50 (2H, s), 3.77 (2H, br), 6.46 (1H, s), 6.55 (1H, d, J = 8.1 Hz), 7.26 (1H, d, J = 8.1 Hz).

Reference Example 71

6-Acetamido-2-tetralone



1) Sodium borohydride (931 mg, 24.6 mmol) was added to a methanol solution (60 ml) of 6-acetamido-1-tetralone (5.00 g, 24.6 mmol) under ice-cooling, which was stirred at room temperature for 1 hour. Ethyl acetate was added to the reaction mixture, which was washed with saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then, the solvent was distilled out under reduced pressure. p-Toluenesulfonic acid (468 mg, 2.46 mmol) and toluene (120 ml) were added to the obtained alcohol form (5.05 g, 24.6 mmol), which was stirred at 100 °C for 1 hour. The solvent was distilled out under reduced pressure. Ethyl acetate was added to the residue, which was washed with saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. The resulting oily substance was purified by silica gel column chromatography (development solvent; hexane:ethyl acetate = 1:1), and powdered with hexane to give N-(7,8-dihydro-2-naphthalenyl)acetamide (3.17 g).

^1H NMR (CDCl_3) δ : 2.16 (3H, s), 2.29 (2H, m), 2.28 (2H, m), 5.97 (1H, m), 6.42 (2H, d, $J=9.6$ Hz), 6.97 (1H, d, $J=8.1$ Hz), 7.14 (1H, br), 7.20 (1H, m), 7.32 (1H, s).

2) m-Chloroperbenzoic acid (5.13 g, 20.8 mmol) was added to a chloroform solution (80 ml) of N-(7,8-dihydro-2-naphthalenyl)acetamide (3.00 g, 16.0 mmol) obtained in 1) under ice-cooling, which was stirred at room temperature for 2 hours. Ethyl acetate was added to the reaction mixture, which was washed with saturated sodium hydrogencarbonate solution and saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure.

The resulting oily substance was purified by alumina B column chromatography (development solvent; hexane:ethyl acetate = 1:1). 1N Sodium hydroxide solution (10.7 ml) was added to a methanol solution (100 ml) of the obtained oily substance (3.20 g, 8.89 mmol) under ice-cooling, which was stirred at room temperature for 30 minutes. The solvent was distilled out under reduced pressure. Ethyl acetate was added to the residue, which was washed with saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. The resulting oily substance was purified by alumina B column chromatography (development solvent; ethyl acetate:methanol = 10:1). p-

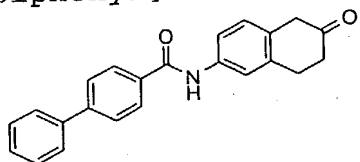
Toluenesulfonic acid (50mg, 0.262 mmol) and toluene (26 ml) were added to the obtained diol (596 mg, 2.62 mmol), which was stirred at 120°C for 3 hours. The solvent was distilled out under reduced pressure. Ethyl acetate was added to the residue, which was washed with saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure.

The resulting oily substance was purified by silica gel column chromatography (development solvent; hexane:ethyl acetate = 1:3), and powdered with diisopropyl ether, to give the titled compound (231 mg).

¹H NMR (CDCl₃) δ: 2.18 (3H, s), 2.54 (2H, m), 3.04 (2H, m), 3.76 (2H, s), 7.06 (1H, d, J=8.1 Hz), 7.21 (1H, dd, J=8.1, 2.0 Hz), 7.31 (1H, br), 7.61 (1H, d, J=2.0 Hz).

5 Reference Example 72

N-(6-Oxo-5,6,7,8-tetrahydro-2-naphthalenyl)[1,1'-biphenyl]-4-carboxamide

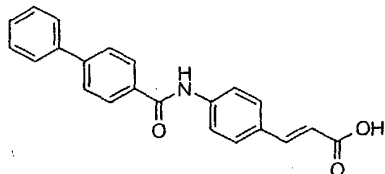


10 Concentrated hydrochloric acid (1.5 ml) was added to 6-acetamido-2-tetralone (20 mg, 0.098 mmol) obtained in Reference Example 71, which was stirred at 100°C for 1 hour, and the solvent was distilled out under reduced pressure. Ethyl acetate was added to the residue, which was washed with aqueous potassium carbonate solution and saturated
15 aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. [1,1'-Biphenyl]-4-carbonyl chloride (21.3 mg, 0.098 mmol) was added to a dimethylformamide solution (0.5 ml) of the obtained oily substance and
20 triethylamine (0.014 ml, 0.098 mmol) under ice-cooling, which was stirred at room temperature for 1 hour. Ethyl acetate was added to the reaction mixture, which was washed with 1N hydrochloric acid, aqueous potassium carbonate solution and saturated aqueous sodium chloride solution,
25 dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. The resulting residue was purified by silica gel column chromatography (development solvent; hexane:ethyl acetate = 1:1), to give the titled compound (10 mg).

30 ¹H NMR (CDCl₃) δ: 2.56 (2H, t, J=6.6 Hz), 3.08 (2H, t, J=6.6 Hz), 3.57 (2H, s), 7.11 (1H, d, J=8.1 Hz), 7.43 (4H, m), 7.64 (2H, m), 7.72 (3H, m), 7.96 (3H, m).

Reference Example 73

(E)-3-[4-[[[1,1'-biphenyl]-4-ylcarbonyl)amino]phenyl]-2-propenic acid

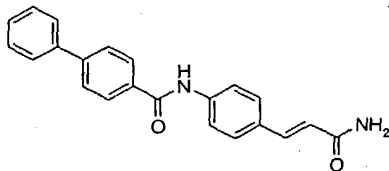


5 4-Phenylbenzoyl chloride (2.00 g, 9.23 mmol) was added to a mixed solution of 4-aminocinnamic acid (1.51 g, 9.23mmol) and sodium hydrogen carbonate (2.33 g, 27.7 mmol) in water and diethyl ether under ice-cooling, which was stirred for 5 hours. After the reaction mixture was
10 separated, 5N hydrochloric acid was added to water layer, and the precipitated crude product was washed with water and ethyl acetate, to give the titled compound (1.34 g). ¹H NMR (DMSO-d₆) δ: 6.84 (1H, d, J = 16.0 Hz), 7.43-7.93 (12H, m), 8.09 (2H, d, J = 8.4 Hz), 10.51 (1H, s).

15

Reference Example 74

N-[4-[(E)-3-Amino-3-oxo-1-propenyl]phenyl][1,1'-biphenyl]-4-carboxamide

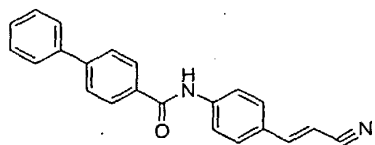


20 Chloro isobutylcarbonate (0.453 ml, 3.49 mmol) was added to a dimethylformamide suspension of (E)-3-[4-[[[1,1'-biphenyl]-4-ylcarbonyl)amino]phenyl]-2-propionic acid (1.00 g, 2.91 mmol) obtained in Reference Example 73 and triethylamine (0.527 ml, 3.79 mmol) under
25 ice-cooling, which was stirred for 30 minute. The solvent was distilled out under reduced pressure. Sodium hydrogencarbonate solution was added to the residue, and the precipitated crude product was washed with water and acetonitrile, to give the titled compound (936 mg).

¹H NMR (DMSO-d₆) δ: 6.56 (1H, d, J = 15.6 Hz), 7.05 (1H, br), 7.52 (7H, m), 7.86 (6H, m), 8.08 (2H, d, J = 7.6 Hz).

Reference Example 75

5 N-[4-[(E)-2-Cyanoethenyl]phenyl][1,1'-biphenyl]-4-carboxamide

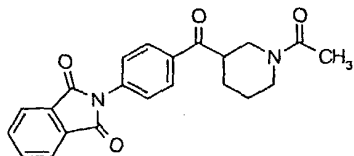


Cyanuric chloride (727 mg, 3.94 mmol) was added to a dimethylformamide suspension of (E)-3-[4-[[[1,1'-
10 biphenyl]-4-ylcarbonyl)amino]phenyl]-2-propenic acid (900 mg, 2.63 mmol) obtained in Reference Example 74 at room temperature, which was stirred for 1 hour. After the solvent was distilled out under reduced pressure, the residue was dissolved in chloroform, which was washed with
15 saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was stillled out under reduced pressure. The resulting residue was purified by silica gel column chromatography (development solvent; chloroform:ethyl acetate = 20:1), to give the
20 titled compound (561 mg) as a colorless powder from diethyl ether.

¹H NMR (DMSO-d₆) δ: 6.37 (1H, d, J = 16.4 Hz), 7.43-7.51 (4H, m), 7.65-7.93 (8H, m), 8.08 (2H, d, J = 8.6 Hz).

25 Reference Example 76

2-[4-[(1-Acetyl-3-piperidiny)carbonyl]phenyl]-1H-isoindol-1,3(2H)-dione



1) Thionyl chloride (2.12 ml, 32.1 mmol) was added to
30 fluorobenzene solution (20 ml) of 1-acetyl-3-

piperidinecarboxylic acid (5.00 g, 29.2 mmol) under ice-cooling, which was stirred at room temperature for 30 minutes. Aluminum chloride (9.74 g, 73.0 mmol) was added to the solution, which was stirred at 90°C for 1 hour. The reaction mixture was poured in ice, and extraction was conducted using ethyl acetate. The extract was washed with saturated aqueous sodium chloride solution, saturated sodium hydrogencarbonate solution, and again saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. The resulting oily substance was purified by silica gel column chromatography (development solvent; hexane:ethyl acetate = 1:1), to give (1-acetyl-3-piperidinyl)(4-fluorophenyl)methanone (4.93 g).
¹H NMR (CDCl₃) δ: 1.61 (2H, m), 1.80 (2H, m), 2.11 and 2.15 (3H, s and s), 2.71 (1H, m), 3.11 and 3.42 (2H, m), 3.87 (1H, m), 4.53 and 4.83 (1H, m), 7.18 (2H, m), 8.02 (2H, m).

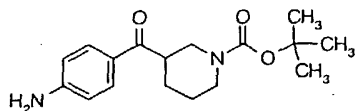
2) A dimethylformamide solution (50 ml) of (1-acetyl-3-piperidinyl)(4-fluorophenyl)methanone (4.92 g, 19.7 mmol) obtained in 1) and potassium phthalimide (3.66g, 19.7mmol) was stirred at 100°C for 12 hours under nitrogen atmosphere. The insoluble matters were filtered off, and the solvent was distilled out under reduced pressure. Ethyl acetate was added to the residue, which was washed with 1N hydrochloric acid and saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure.

The resulting oily substance was purified by silica gel column chromatography (development solvent; ethyl acetate), to give the titled compound (4.18 g) as a colorless powder from ethyl acetate - diisopropyl ether (1:5).

¹H NMR (CDCl₃) δ: 1.66 (2H, m), 1.86 (2H, m), 2.13 and 2.15 (3H, s and s), 2.74 (1H, m), 3.11 and 3.43 (2H, m), 3.88 (1H, m), 4.54 and 4.85 (1H, m), 7.66 (2H, m), 7.82 (2H, m), 7.99 (2H, m), 8.10 (2H, m).

Reference Example 77

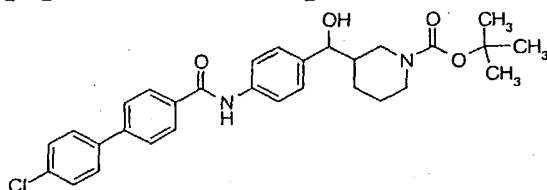
tert-Butyl 3-(4-aminobenzoyl)-1-piperidinecarboxylate



- 5 1) Concentrated hydrochloric acid (53 ml) was added to 2-[4-[(1-acetyl-3-piperidinyl)carbonyl]phenyl]-1H-isoindol-1,3(2H)-dione (4.00 g, 10.6 mmol) obtained in Reference Example 76, which was stirred at 100°C for 16 hours, and then insoluble matters were filtered off.
- 10 Potassium carbonate was added to the filtrate to make it alkaline, and extraction was conducted using ethyl acetate. The extract was washed with saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure.
- 15 The resulting residue was powdered with diisopropyl ether, to give (4-aminophenyl)(3-piperidinyl)methanone (1.69 g). ¹H NMR (CD₃OD) δ : 1.59-1.85 (4H, m), 2.68-2.72 (2H, m), 3.30 (2H, m), 3.45 (1H, m), 6.62 (2H, m), 7.74 (2H, m).
- 20 2) t-Butyl dicarbonate (0.562 ml, 2.45 mmol) was added to a tetrahydrofuran solution (12 ml) of (4-aminophenyl)(3-piperidinyl)methanone (500 mg, 2.45 mmol) obtained in 1) under ice-cooling, which was stirred for 1.5 hours. Ethyl acetate was added to the reaction mixture, which was washed with saturated sodium hydrogencarbonate solution and saturated aqueous sodium chloride solution,
- 25 dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. The resulting oily substance was purified by silica gel column chromatography (development solvent; hexane:ethyl acetate = 1:1), to give the titled compound (831 mg).
- 30 ¹H NMR (CDCl₃) δ 1.47 (9H, s), 1.47-1.52 (2H, m), 1.67-1.74 (2H, m), 2.00 (1H, m), 2.72 (1H, m), 2.90 (1H, m), 3.32 (1H, m), 4.13 (3H, m), 6.66 (2H, d, J=8.4Hz), 7.84 (2H, d, J=8.4Hz).

Reference Example 78

tert-Butyl 3-[[4-[[[(4'-chloro[1,1'-biphenyl]-4-yl)carbonyl]amino]phenyl](hydroxy)methyl]-1-piperidinecarboxylate

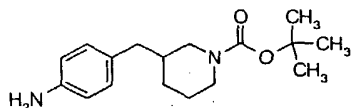


tert-Butyl 3-[4-[[[(4'-chloro[1,1'-biphenyl]-4-yl)carbonyl]amino]benzoyl]-1-piperidinecarboxylate (506 mg, 0.975 mmol) obtained in Example 127-1) was dissolved in a mixed solution of methanol and tetrahydrofuran (1:1) (10 ml). Sodium borohydride (73.8 mg, 1.95 mmol) was added to the solution under ice-cooling, which was stirred at room temperature for 1 hour. Ethyl acetate was added to the reaction mixture, which was washed with saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. Diisopropyl ether was added to the residue, to give the titled compound (488mg) as a colorless powder.

FABMS(pos) 521.2 [M+H]⁺

Reference Example 79

tert-Butyl 3-(4-aminobenzyl)-1-piperidinecarboxylate



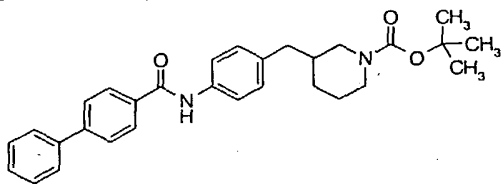
Sodium borohydride (433 mg, 11.5 mmol) was added to a methanol solution (25 ml) of tert-butyl 3-(4-aminobenzoyl)-1-piperidinecarboxylate (1.74g, 5.73mmol) obtained in Reference Example 77 under ice-cooling, which was stirred at room temperature for 1 hour. Ethyl acetate was added to the reaction mixture, which was washed with saturated aqueous sodium chloride solution, dried over

anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. The resulting oily substance was purified by alumina B column chromatography (development solvent; ethyl acetate), to give an alcohol form. 1N hydrochloric acid (9.79 ml) and 10% palladium carbon (200 mg) were added to a methanol solution (300 ml) of the obtained alcohol form (1.00 g, 3.26 mmol), which was stirred for 16 hours under hydrogen atmosphere. The catalyst was filtered off, potassium carbonate was added to the filtrate to make it alkaline, and then the solvent was distilled out under reduced pressure. Ethyl acetate was added to the residue, which was washed with saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. The resulting oily substance was purified by silica gel column chromatography (development solvent; hexane - ethyl acetate = 1:1), to give the titled compound (813 mg).

^1H NMR (CDCl_3) δ : 1.46-1.76 (14H, m), 2.25-2.80 (2H, m), 3.14 (2H, m), 3.76 (4H, m), 6.64 (2H, m), 7.01 (2H, m).

Reference Example 80

tert-Butyl 3-[4-[[[1,1'-biphenyl]-4-ylcarbonyl)amino]benzyl]-1-piperidinecarboxylate



25

The titled compound was obtained by carrying out the same operation as in Example 1, using tert-butyl 3-(4-aminobenzyl)-1-piperidinecarboxylate obtained in Reference Example 79 and [1,1'-biphenyl]-4-carboxylic acid.

30

Elemental analysis for $\text{C}_{30}\text{H}_{34}\text{N}_2\text{O}_3 \cdot 0.5\text{H}_2\text{O}$

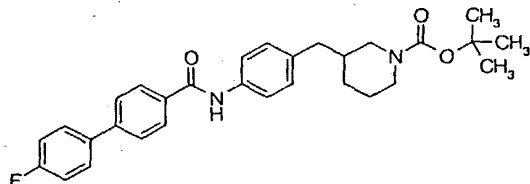
Calcd.: C, 75.13; H, 7.36; N, 5.84.

Found: C, 74.83; H, 7.25; N, 5.65.

Melting point: 135 - 137°C

Reference Example 81

- 5 tert-Butyl 3-[4-[[[(4'-fluoro[1,1'-biphenyl]-4-yl)carbonyl]amino]benzyl]-1-piperidinecarboxylate



- The titled compound was obtained by carrying out the same operation as in Example 1, using tert-butyl 3-(4-aminobenzyl)-1-piperidinecarboxylate obtained in Reference Example 80 and 4'-fluoro[1,1'-biphenyl]-4-carboxylic acid.

Elemental analysis for $C_{30}H_{33}FN_2O_3 \cdot 0.5H_2O$

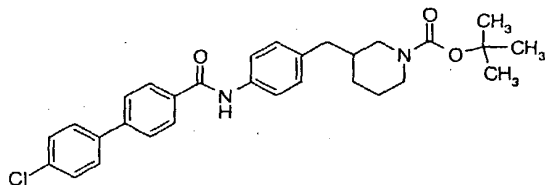
Calcd.: C, 72.41; H, 6.89; N, 5.63.

- 15 Found: C, 72.30; H, 7.07; N, 5.60.

Melting point: 138 - 141°C

Reference Example 82

- 20 tert-Butyl 3-[4-[[[(4'-chloro[1,1'-biphenyl]-4-yl)carbonyl]amino]benzyl]-1-piperidinecarboxylate



- The titled compound was obtained by carrying out the same operation as in Example 1, using tert-butyl 3-(4-aminobenzyl)-1-piperidinecarboxylate obtained in Reference Example 80 and 4'-chloro[1,1'-biphenyl]-4-carboxylic acid.

Elemental analysis for $C_{30}H_{33}ClN_2O_3 \cdot 0.5H_2O$

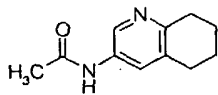
Calcd.: C, 70.09; H, 6.67; N, 5.45.

Found: C, 70.29; H, 6.50; N, 5.38.

Melting point: 173 - 176°C

Reference Example 83

N-(5,6,7,8-Tetrahydro-3-quinoliny)acetamide



1) Fuming nitric acid (100 ml) was added dropwise to concentrated sulfuric acid solution (200 ml) of 1-methyl-2-pyridone (20.7 g, 190 mmol) at 100°C, which was stirred for 16 hours. The reaction mixture was poured in ice. The resulting precipitate was collected, which was washed with water, to give 1-methyl-3,5-dinitro-2(1H)-pyridinone (3.0 g).

¹H NMR (DMSO-d₆) δ: 3.68 (3H, s), 9.01 (1H, d, J=3.0 Hz), 9.61 (1H, d, J=3.0 Hz).

15 2) 1N Methanolic ammonia solution (300 ml) of 1-methyl-3,5-dinitro-2(1H)-pyridinone (3.00g, 15.1mmol) obtained in 1) and 1-morpholino-1-cyclohexene (3.88 ml, 22.6 mmol) was stirred at 70°C for 3 hours. The solvent was distilled out under reduced pressure. The resulting residue was purified by alumina column chromatography (development solvent; ethyl acetate), to give 3-nitro-5,6,7,8-tetrahydroquinoline (2.42 g) as a powder from methanol - water (1:4).

25 ¹H NMR (DMSO-d₆) δ: 1.87 (4H, m), 2.90 (4H, m), 8.15 (1H, s), 9.16 (1H, s).

3) 10% Palladium-carbon (200 mg) was added to a methanol solution (68 ml) of 3-nitro-5,6,7,8-tetrahydroquinoline (2.41 g, 13.5 mmol) obtained in 2), which was stirred under hydrogen atmosphere for 16 hours.

30 After a catalyst was filtered off, the solvent was distilled out under reduced pressure. The resulting residue was dissolved in pyridine (35 ml). Anhydrous ethyl acetate (1.91 ml, 20.3 mmol) was added to the solution, which was stirred at room temperature for 1 hour. After completion of the reaction, the solvent was distilled out

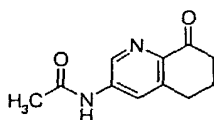
35

under reduced pressure. Diisopropyl ether - n-hexane (1:8) was added to the resulting residue, to give the titled compound (2.48 g) as a colorless powder.

¹H NMR (CDCl₃) δ: 1.80-1.87 (4H, m), 2.18 (3H, s), 2.77 (2H, m), 2.87 (2H, m), 7.72 (1H, br), 7.94 (1H, s), 8.24 (1H, s).

Reference Example 84

N-(8-Oxo-5,6,7,8-tetrahydro-3-quinolinyl)acetamide



10

1) m-Chloroperbenzoic acid (3.83 g, 15.5 mmol) was added to a chloroform solution (65 ml) of N-(5,6,7,8-tetrahydro-3-quinolinyl)acetamide (2.46 g, 12.9 mmol) obtained in Reference Example 83 under ice-cooling, which was stirred at room temperature for 16 hours. After the solvent was distilled out under reduced pressure, the residue was powdered with ethyl acetate, to give N-(1-oxide-5,6,7,8-tetrahydro-3-quinolinyl)acetamide (2.00 g).

20 ¹H NMR (DMSO-d₆) δ: 1.64 (2H, m), 1.75 (2H, m), 2.04 (3H, s), 2.66 (4H, m), 7.13 (1H, s), 8.56 (1H, s), 10.12 (1H, s).

2) Anhydrous ethyl acetate (30 ml) was added to N-(1-oxide-5,6,7,8-tetrahydro-3-quinolinyl)acetamide (1.99 g, 9.65 mmol) obtained in 1), which was stirred at 80°C for 3 hours. The reaction mixture was cooled to room temperature. The solvent was distilled out under reduced pressure, and the resulting residue was purified by alumina column chromatography (development solvent; ethyl acetate). The resulting oily substance was dissolved in methanol (110 ml). 1N Sodium hydroxide (21.5 ml) was added to the solution under ice-cooling, which was stirred at room temperature for 1 hour. The solvent was distilled out under reduced pressure. Chloroform was added to the residue,

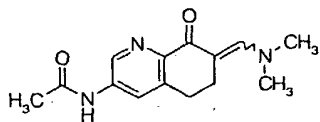
30

which was washed with aqueous potassium carbonate solution and saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. The resulting
5 residue was purified by alumina column chromatography (development solvent; ethyl acetate:methanol = 5:1), to give N-(8-hydroxy-5,6,7,8-tetrahydro-3-quinolinyl)acetamide (1.08 g) as a powder from ethyl acetate and diisopropyl ether.
10 ¹H NMR (CDCl₃) δ: 1.79 (2H, m), 1.96 (1H, m), 2.22 (3H, s), 2.24 (1H, m), 2.82 (2H, m), 4.69 (1H, m), 7.49 (1H, br), 7.92 (1H, s), 8.30 (1H, s).

3) Manganese dioxide (4.47 g, 51.4 mmol) was added to chloroform (26 ml) solution of N-(8-hydroxy-5,6,7,8-
15 tetrahydro-3-quinolinyl)acetamide (1.06 g, 5.14 mmol) obtained in 2), which was stirred at room temperature for 1 day. After completion of the reaction, the insoluble matters were filtered off, and the filtrate was concentrated under reduced pressure. Diisopropyl ether
20 and hexane were added to the resulting residue, to give the titled compound (858 mg) as a colorless powder.
¹H NMR (CDCl₃) δ: 2.20 (2H, m), 2.26 (3H, s), 2.77 (2H, m), 3.03 (2H, m), 8.10 (1H, br), 8.39 (1H, s), 8.42 (1H, s).

25 Reference Example 85

N-[7-[(Dimethylamino)methylidene]-8-oxo-5,6,7,8-tetrahydro-3-quinolinyl]acetamide

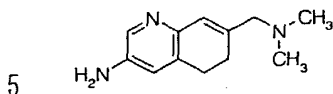


The titled compound was obtained by carrying out the
30 same operation as in Reference Example 47, using N-(8-oxo-5,6,7,8-tetrahydro-3-quinolinyl)acetamide obtained in Reference Example 84.

¹H NMR (CDCl₃) δ: 2.09 (3H, s), 2.78 (2H, m), 2.85 (2H, m), 3.10 (6H, s), 7.55 (1H, s), 8.01 (1H, s), 8.56 (1H, s).

Reference Example 86

N-[(3-Amino-5,6-dihydro-7-quinolinyl)methyl]-N,N-dimethylamine



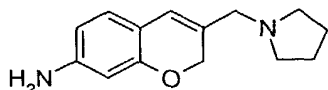
The titled compound was obtained by carrying out the same operation as in Reference Example 41-2), using N-[7-[(dimethylamino)methylidene]-8-oxo-5,6,7,8-tetrahydro-3-quinolinyl]acetamide obtained in Reference

10 Example 85.

¹H NMR (CDCl₃) δ: 2.23 (6H, s), 2.33 (2H, t, J=8.1 Hz), 2.78 (2H, t, J=8.1 Hz), 2.99 (2H, s), 3.59 (2H, br), 6.43 (1H, s), 6.74 (1H, d, J=2.5 Hz), 7.84 (1H, d, J=2.5 Hz).

15 Reference Example 87

3-(1-Pyrrolidinylmethyl)-2H-chromen-7-amine



The titled compound was obtained as an oily substance by carrying out the same operations as in Example 41-1),

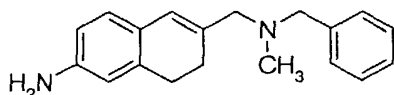
20 Reference Example 52 and Example 41-2) in this order, using 7-acetylamino-3,4-dihydrochromen-4-one.

¹H-NMR (CDCl₃) δ: 1.77-1.79 (4H, m), 2.45-2.47 (4H, m), 3.11 (2H, s), 3.66 (2H, s), 4.74 (2H, s), 6.14-6.21 (3H, m), 6.75 (1H, d, J = 7.8 Hz).

25

Reference Example 88

6-[(N-Benzyl-N-methylamino)methyl]-7,8-dihydro-2-naphthalenamine



30

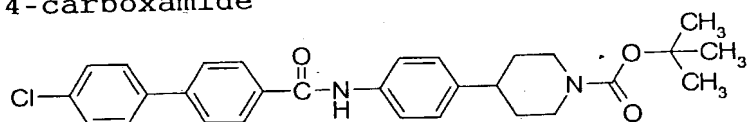
The titled compound was obtained as an oily substance by carrying out the same operation as in Reference Example 52, using 6-acetamido-2-(N,N-dimethylaminomethylidene)-

1-tetralone obtained in Example 41-1).

¹H-NMR (CDCl₃) δ: 2.17 (3H, s), 2.35 (2H, t, J = 8.1 Hz),
2.73 (2H, t, J = 8.1 Hz), 3.04 (2H, s), 3.48 (2H, s), 3.58
(2H, s), 6.29 (1H, s), 6.44 - 6.46 (2H, m), 6.82 (1H, d, J
5 = 8.1 Hz), 7.03-7.45 (5H, m).

Reference Example 89

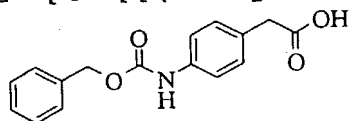
4'-Chloro-N-[4-(4-piperidiny)phenyl][1,1'-biphenyl]-
4-carboxamide



10 An ethanol solution (30 ml) of tert-butyl 4-(4-nitrophenyl)-1-piperidinecarboxylate (1.7 g) was subjected to catalytic hydrogenation using 10% palladium carbon (0.2 g) as a catalyst under normal temperature and
15 normal pressure. After the catalyst was filtered off, the filtrate was concentrated to give tert-butyl 4-(4-aminophenyl)-1-piperidinecarboxylate as a viscous oily substance. The titled compound (2.2 g) was obtained as
20 colorless crystals, by carrying out the same operation as in Example 1, using the resulting oily substance and 4'-chloro[1,1'-biphenyl]-4-carboxylic acid (1.43 g).
¹H-NMR (CDCl₃+ DMSO-d₆) δ: 1.05-1.32 (11H, m), 1.38-1.50 (2H, m), 2.20-2.50 (3H, m), 3.75-3.90 (2H, m), 6.81 (2H, d, J=8.4 Hz), 7.07 (2H, d, J=8.4 Hz), 7.20-7.36 (6H, m),
25 7.69 (2H, d, J=8.1Hz), 9.44 (1H, s).
Melting point: 232 - 233°C (crystallization solvent : ethyl acetate)

Reference Example 90

30 2-[4-[(Benzyloxy)carbonyl]amino]phenyl]ethyl acetate

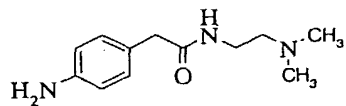


To an ethyl acetate (100 ml) suspension of 4-

aminophenylethyl acetate (10 g), saturated aqueous sodium bicarbonate solution (100 ml) was added, and further, benzyloxycarbonyl chloride (12.3 ml) was added dropwise under ice-cooling. After stirring for 1 hour,
 5 hydrochloric acid was added to the reaction mixture to make it acidic, and extraction was conducted using ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was recrystallized from ethyl
 10 acetate - hexane, to give the titled compound (17.3 g). Melting point: 148 - 149°C

Reference Example 91

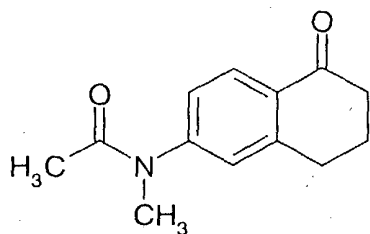
2-(4-Aminophenyl)-N-[2-(
 15 (dimethylamino)ethyl]acetamide



Pd-C (1 g) was added to a methanol (140 ml) solution of benzyl 4-[2-[[2-(dimethylamino)ethyl]amino]-2-oxoethyl]phenylcarbamate (10 g), which was stirred under
 20 hydrogen atmosphere for 1 hour. Pd-C was removed, and the filtrate was concentrated. The residue was purified by alumina column chromatography (development solvent; ethyl acetate:hexane = 1:1), to give the titled compound (6.63 g) as an oily substance.
 25 ¹H-NMR(CDCl₃) δ: 2.16 (6H, s), 2.05 (3H, s), 2.30-2.36 (2H, t, J=6.2 Hz), 3.23-3.32 (2H, dd, J=11.4, 6.2 Hz), 3.44 (2H, s), 6.00 (1H, s), 6.63-6.67 (2H, m), 7.00-7.07 (2H, m).

30 Reference Example 92

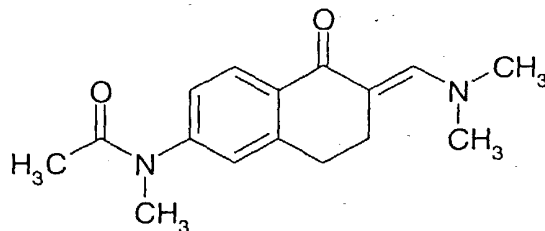
N-Methyl-N-(5-oxo-5,6,7,8-tetrahydro-2-naphthalenyl)acetamide



6-Acetamido-1-tetralone (10.0 g, 49.2 mmol) was dissolved in tetrahydrofuran (100 ml). Sodium hydride (oil, 3.0 g) was added to the solution, which was refluxed with heating under nitrogen atmosphere for 2 hours. After cooling, methyl iodide (30 ml) was added to the reaction mixture, which was refluxed with heating under nitrogen atmosphere for 2 hours. The reaction mixture was concentrated. Ethyl acetate and water were added to the residue, and extraction was conducted. The ethyl acetate layer was concentrated, and the residue was purified by alumina column chromatography (development solvent; ethyl acetate:n-hexane = 33:67 ~ 50:50). The product was concentrated under reduced pressure, and the residue was recrystallized from ethyl acetate - diisopropyl ether, to give the titled compound (4.3 g).
¹H-NMR (CDCl₃) δ: 1.96 (3H, brs), 2.18 (2H, m), 2.69 (2H, t, J=6.1 Hz), 2.99 (2H, t, J=5.9 Hz), 3.29 (3H, s), 7.01-7.15 (2H, m), 8.08 (1H, d, J=8.1 Hz).

Reference Example 93

N-[6-[(Dimethylamino)methylidene]-5-oxo-5,6,7,8-tetrahydro-2-naphthalenyl]-N-methylacetamide



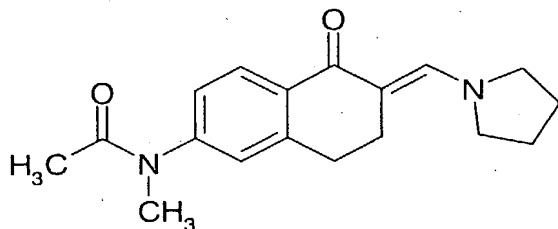
N-Methyl-N-(5-oxo-5,6,7,8-tetrahydro-2-naphthalenyl)acetamide (4.3 g, 19.8 mmol) obtained in Reference Example 92 was dissolved in N,N-

dimethylformamide dimethylacetal (50 ml), which was refluxed with heating under nitrogen atmosphere for 15 hours. The reaction mixture was concentrated, and the residue was washed with ethyl acetate and diisopropyl ether, to give the titled compound (3.9 g).

$^1\text{H-NMR}$ (CDCl_3) δ : 1.93 (3H, brs), 2.84 (2H, dd, $J=7.5$, 5.6 Hz), 2.95 (2H, dd, $J=7.5$, 5.6 Hz), 3.16 (6H, s), 3.28 (3H, s), 6.99 (1H, s), 7.10 (1H, dd, $J=8.1$, 2.0 Hz), 7.75 (1H, s), 8.07 (1H, d, $J=8.1$ Hz).

Reference Example 94

N-Methyl-N-[5-oxo-6-[1-pyrrolidinylmethylidene]-5,6,7,8-tetrahydro-2-naphthalenyl]acetamide



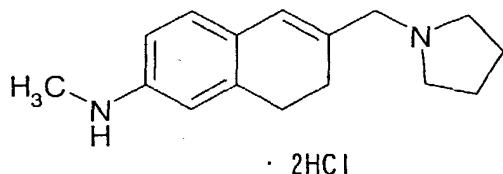
N-[6-[(Dimethylamino)methylidene]-5-oxo-5,6,7,8-tetrahydro-2-naphthalenyl]-N-methylacetamide (5.7 g, 20.9 mmol) obtained in Reference Example 93 was dissolved in pyrrolidine (50 ml), which was refluxed with heating under nitrogen atmosphere for 3.5 hours. Then, ethyl acetate and water were added to the reaction mixture, and extraction was conducted. The ethyl acetate layer was concentrated, and the residue was recrystallized from ethyl acetate - diisopropyl ether, to give the titled compound (4.0 g, yield : 64%).

$^1\text{H-NMR}$ (CDCl_3) δ : 1.94 (7H, m), 2.84 (2H, dd, $J=7.0$, 5.6 Hz), 2.97 (2H, dd, $J=7.0$, 5.6 Hz), 3.28 (3H, s), 3.63 (4H, m), 6.98 (1H, s), 7.10 (1H, dd, $J=8.1$, 2.0 Hz), 7.95 (1H, s), 8.08 (1H, d, $J=8.1$ Hz).

Reference Example 95

N-Methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-

naphthalenamine dihydrochloride

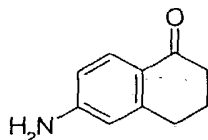


N-Methyl-N-[5-oxo-6-[1-pyrrolidinylmethylidene]-5,6,7,8-tetrahydro-2-naphthalenyl]acetamide (4.0 g, 13.4 mmol) obtained in Reference Example 94 was dissolved in methanol - ethyl acetate (10:1, 220 ml). 10% Palladium carbon (50% wet, 0.4 g) was added to the solution, which was ice cooled. Stirring was begun under hydrogen atmosphere, and stirring was conducted for 2 days while returning the temperature of the reaction mixture to room temperature. A catalyst was filtered off, the reaction mixture was concentrated under reduced pressure, and the residue was dissolved in ethyl acetate. Extraction was conducted using 1N hydrochloric acid. The extract was made alkaline with 4N sodium hydroxide solution, and extraction was conducted using ethyl acetate. The extract was concentrated under reduced pressure. The residue was dissolved in tetrahydrofuran (100 ml) and 5N hydrochloric acid (100 ml), which was refluxed with heating for 13 hours. The reaction mixture was concentrated. Ethyl acetate and saturated aqueous sodium carbonate solution were added to the residue, and extraction was conducted. The ethyl acetate layer was concentrated. 4N Hydrogen chloride - ethyl acetate solution was added to the resulting oily substance, which was concentrated. The residue was recrystallized from methanol - ethyl acetate, to give the titled compound (2.8 g, yield : 66%).

¹H-NMR (DMSO-d₆) δ: 1.98 (4H, m), 2.45 (4H, m), 2.81 (5H, m), 3.01 (2H, brd), 3.44 (2H, brd), 3.86 (2H, d, J=5.0 Hz), 7.02-7.10 (3H, m), 10.89 (1H, brs).

Reference Example 96

6-Amino-3,4-dihydro-1-(2H)-naphthalenone

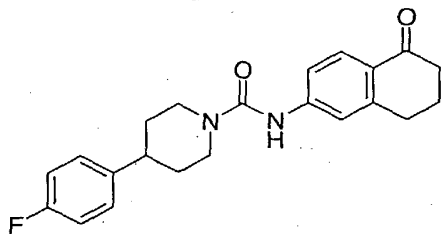


Concentrated hydrochloric acid (250 ml) was added to 6-acetamido-1-tetralone (20.0 g, 98.4 mmol), which was stirred at 100°C for 1 hour. The solvent was distilled out under reduced pressure. Ethyl acetate was added to the residue, which was washed with aqueous potassium carbonate solution and saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. The residue was powdered with ethyl acetate and isopropyl ether, to give the titled compound (14.5 g).

¹H NMR (CDCl₃) δ: 2.07 (2H, m), 2.57 (2H, m), 2.83 (2H, m), 4.10 (2H, br), 6.42 (1H, d, J=2.2 Hz), 6.53 (1H, dd, J=2.2, 8.4Hz), 7.89 (1H, d, J=8.4 Hz).

Reference Example 97

4-(4-Fluorophenyl)-N-(5-oxo-5,6,7,8-tetrahydro-2-naphthalenyl)-1-piperidinecarboxamide



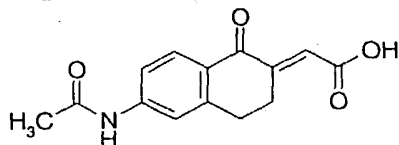
Pyridine(9.95 ml, 123 mmol) and 4-nitrophenyl chloroformate (12.4 g, 61.5 mmol) was added to a tetrahydrofuran(300 ml)solution of 6-amino-3,4-dihydro-1(2H)-naphthalenone(9.92 g, 61.5 mmol)obtained in Reference Example 96, which was stirred at room temperature for 3 hours. The solvent was distilled out under reduced pressure. 1N Hydrochloric acid was added to the residue to powder, which was washed with ethanol. 4N Aqueous sodium hydroxide solution was added to a dimethylsulfoxide (33 ml)solution of the resulting 4-nitrophenyl-5-oxo-

5,6,7,8-tetrahydro-2-naphthalenylcarbamate (2.20 g, 6.74 mmol) and 4-(4-fluorophenyl)piperidine hydrochloride (1.60 g, 7.42 mmol), which was stirred at room temperature for 1 hour. Ethyl acetate was added to the reaction mixture, which was washed with 1N hydrochloric acid, aqueous potassium hydrogencarbonate solution and saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and the solvent was distilled out under reduced pressure. The resulting residue was purified by alumina B column chromatography (development solvent; ethyl acetate), and powdered with isopropyl ether and hexane, to give the titled compound (1.89 g).

¹H NMR (CDCl₃) δ: 1.72 (2H, m), 1.92 (2H, m), 2.11 (2H, m), 2.61 (2H, m), 2.72 (1H, m), 2.93 (2H, m), 3.01 (2H, m), 4.23 (2H, m), 6.67 (1H, s), 7.00 (2H, m), 7.12 (3H, m), 7.61 (1H, s), 7.97 (1H, d, J=8.4 Hz).

Reference Example 98

[6-(Acetylamino)-1-oxo-3,4-dihydro-2(1H)-naphthalenyldene]acetic acid

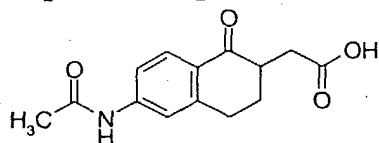


0.5N Aqueous sodium hydroxide solution (190 ml) was added to an aqueous solution (60 ml) of 6-acetamido-1-tetralone (5.00 g, 24.6 mmol) and glyoxylic acid (9.05 g, 98.5 mmol) under ice-cooling, which was stirred at 60°C for 16 hours. After cooling, concentrated hydrochloric acid was added to the reaction mixture. The precipitated crystals were collected, which was washed with water, to give the titled compound (3.73 g).

¹H NMR (DMSO-d₆) δ: 2.10 (3H, s), 2.95 (2H, m), 3.28 (2H, m), 6.63 (1H, s), 7.53 (1H, d, J=8.7Hz), 7.67 (1H, s), 7.91 (1H, d, J=8.7Hz), 10.32 (1H, s), 12.89 (1H, br).

Reference Example 99

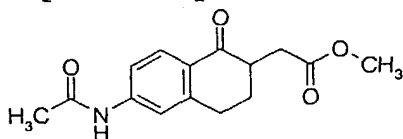
[6-(Acetylamino)-1-oxo-1,2,3,4-tetrahydro-2-naphthalenyl]acetic acid



5 70% Acetic acid - water solution (35 ml) of [6-(acetylamino)-1-oxo-3,4-dihydro-2(1H)-naphthalenyliden]acetic acid (3.50 g, 13.5 mmol) obtained in Reference Example 98 and zinc powder (2.1 g) was stirred at 100°C for 30 minutes. After cooling, zinc powder was
10 filtered. Ethyl acetate was added to the filtrate, which was washed with saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. The resulting oily substance was purified by silica gel column
15 chromatography (development solvent; ethyl acetate : methanol = 10:1), and powdered with ethyl acetate and isopropyl ether, to give the titled compound (2.51 g).
¹H NMR (CDCl₃) δ : 1.85-2.15 (2H, m), 2.08 (3H, s), 2.38 (1H, m), 2.71 (1H, m), 2.88 (2H, m), 3.05 (1H, m), 7.46 (1H, d, J=8.7Hz), 7.60 (1H, s), 7.80 (1H, d, J=8.7Hz), 10.21 (1H, s), 12.09 (1H, br).

Reference Example 100

25 Methyl [6-(acetylamino)-1-oxo-1,2,3,4-tetrahydro-2-naphthalenyl]acetate



Methyl iodide (0.18 ml, 2.87 mmol) was added to a dimethylformamide solution (10 ml) of [6-(acetylamino)-1-oxo-1,2,3,4-tetrahydro-2-naphthalenyl]acetic acid (500
30 mg, 1.91 mmol) obtained in Reference Example 99 and potassium carbonate (529 mg, 3.82 mmol), which was stirred

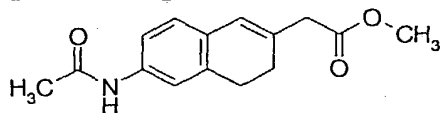
at room temperature for 16 hours. Ethyl acetate was added to the reaction mixture, which was washed with aqueous sodium thiosulfate solution and saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure.

The resulting oily substance was purified by alumina B column chromatography (development solvent; ethyl acetate), to give the titled compound (527 mg).

¹H NMR (CDCl₃) δ: 1.98 (1H, m), 2.20 (3H, s), 2.23 (1H, m), 2.47 (1H, m), 3.30 (4H, m), 3.73 (3H, s), 7.21 (1H, d, J=8.7Hz), 7.50-7.80 (2H, m), 7.97 (1H, d, J=8.7Hz).

Reference Example 101

Methyl [6-(acetylamino)-3,4-dihydro-2-naphthalenyl]acetate



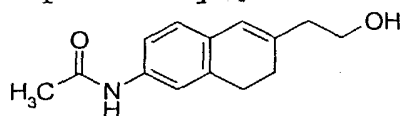
Sodium borohydride (72.4 mg, 1.91 mmol) was added to a methanol solution (10ml) of methyl [6-(acetylamino)-1-oxo-1,2,3,4-tetrahydro-2-naphthalenyl]acetate (527 mg, 1.91 mmol) obtained in Reference Example 100 under ice-cooling, which was stirred for 1 hour. Ethyl acetate was added to the reaction mixture, which was washed with saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. The resulting oily substance was purified by alumina B column chromatography (development solvent; ethyl acetate). Concentrated sulfuric acid (0.14 ml) was added to an acetic acid solution (7 ml) of the oil (404 mg, 1.46 mmol), which was stirred at 40°C for 5 hours. The solvent was distilled out under reduced pressure. Ethyl acetate was added to the residue, which was washed with aqueous potassium carbonate solution and saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was

distilled out under reduced pressure. The resulting oily substance was purified by silica gel column chromatography (development solvent; hexane:ethyl acetate = 1:1), to give the titled compound (251 mg).

- 5 ^1H NMR (CDCl_3) δ : 2.16 (3H, s), 2.32 (2H, t, $J=8.1\text{Hz}$), 2.82 (2H, t, $J=8.1\text{Hz}$), 3.21 (2H, s), 3.71 (3H, s), 6.30 (1H, s), 6.93 (1H, d, $J=8.1\text{Hz}$), 7.19 (2H, m), 7.33 (1H, s).

Reference Example 102

- 10 N-[6-(2-Hydroxyethyl)-7,8-dihydro-2-naphthalenyl]acetamide

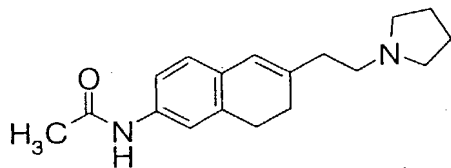


- Lithium aluminum hydride (242 mg, 6.38 mmol) was added to a tetrahydrofuran solution (16 ml) of methyl [6-(acetylamino)-3,4-dihydro-2-naphthalenyl]acetate (827 mg, 3.19 mmol) obtained in Reference Example 101 under ice-cooling, which was stirred at room temperature for 1 hour. Ethyl acetate was added to the reaction mixture, which was washed with 1N hydrochloric acid and saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. The residue was powdered with isopropyl ether, to give the titled compound (364 mg).
- 15
- 20

- 25 ^1H NMR (CDCl_3) δ : 1.43 (1H, m), 2.16 (3H, s), 2.26 (2H, t, $J=8.1\text{Hz}$), 2.46 (2H, t, $J=6.3\text{Hz}$), 2.81 (2H, t, $J=8.1\text{Hz}$), 3.78 (2H, m), 6.28 (1H, s), 6.94 (1H, d, $J=8.1\text{Hz}$), 7.08 (1H, br), 7.17 (1H, d, $J=8.1\text{Hz}$), 7.35 (1H, s).

Reference Example 103

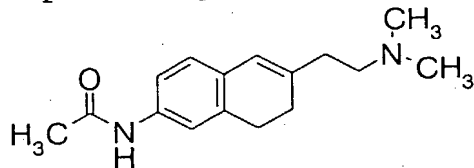
- 30 N-[6-[2-(1-Pyrrolidinyl)ethyl]-7,8-dihydro-2-naphthalenyl]acetamide



Methanesulfonyl chloride (0.131 ml, 1.69 mmol) was added to a dimethylformamide solution (7 ml) of N-[6-(2-hydroxyethyl)-7,8-dihydro-2-naphthalenyl]acetamide (355 mg, 1.53 mmol) obtained in Reference Example 102 and triethylamine (0.235 ml, 1.69 mmol) under ice-cooling, which was stirred for 30 minutes. Pyrrolidine (0.384 ml, 4.60 mmol) was added to the reaction mixture, which was stirred at 60 °C for 4 hours. The solvent was distilled out under reduced pressure. Ethyl acetate was added to the residue, and extraction was conducted using 1N hydrochloric acid. Potassium carbonate was added to the extract to make it alkaline, and extraction was conducted using ethyl acetate. The extract was washed with saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and the solvent was distilled out under reduced pressure. The resulting residue was purified by alumina column chromatography (development solvent; ethyl acetate), to give the titled compound (294 mg).

¹H NMR (CDCl₃) δ: 1.79 (4H, m), 2.16 (3H, s), 2.25 (2H, m), 2.41 (2H, m), 2.55 (4H, m), 2.62 (2H, m), 2.78 (2H, m), 6.20 (1H, s), 6.91 (1H, d, J=8.1Hz), 7.18 (1H, d, J=7.8Hz), 7.32 (2H, m).

Reference Example 104
N-[6-[2-(Dimethylamino)ethyl]-7,8-dihydro-2-naphthalenyl]acetamide



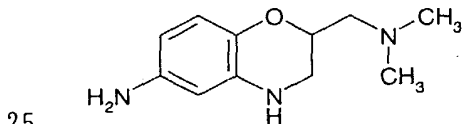
Methanesulfonyl chloride (0.0393 ml, 0.469 mmol) was added to a dimethylformamide solution (2 ml) of N-[6-

(2-hydroxyethyl)-7,8-dihydro-2-naphthalenyl]acetamide (102 mg, 0.426 mmol) obtained in Reference Example 102 and triethylamine (0.0652 ml, 0.469 mmol) under ice-cooling, which was stirred for 30 minutes. A tetrahydrofuran solution (0.64 ml) of 2N dimethylamine was added to the reaction mixture, which was stirred at 60°C for 5 hours. The solvent was distilled out under reduced pressure. Ethyl acetate was added to the residue, and extraction was conducted using 1N hydrochloric acid. Potassium carbonate was added to the extract to make it alkaline, and extraction was conducted using ethyl acetate. The extract was washed with saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and the solvent was distilled out under reduced pressure. The resulting residue was purified by alumina column chromatography (development solvent; ethyl acetate), to give the titled compound (57.5 mg).

¹H NMR (CDCl₃) δ: 2.15 (3H, s), 2.24 (2H, m), 2.29 (6H, s), 2.36 (2H, m), 2.48 (2H, m), 2.78 (2H, m), 6.20 (1H, s), 6.90 (1H, d, J=8.1Hz), 7.20 (1H, d, J=8.1Hz), 7.35 (1H, s), 7.76 (1H, br).

Reference Example 105

6-Amino-2-[(dimethylamino)methyl]-1,4-benzoxazine



1) 2-Ethoxycarbonyl-6-nitro-1,4-benzoxazine (7.20 g, 0.029 mol) obtained by a known method by documents (Journal of heterocyclic chemistry, 19(5), p.1189 (1982)) was dissolved in methanol (50 ml). Sodium borohydride (1.08 g, 0.029 mol) was added to the solution, which was stirred for 2 hours. The reaction mixture was concentrated. Ethyl acetate and aqueous potassium hydrogencarbonate solution were added to the residue, and extraction was conducted. The organic layer was washed

with water, and concentrated. A mixed solution of ethyl acetate and n-hexane (1:5) was added to the residue for crystallization. The crystallized product was collected by filtration, to give 2-hydroxymethyl-6-nitro-1,4-benzoxazine (3.10 g) as a red powder.

$^1\text{H-NMR}$ (CDCl_3) δ : 1.96 (1H, m), 3.34-3.49 (2H, m), 3.80-3.90 (2H, m), 4.09 (1H, brs), 4.30-4.40 (1H, m), 6.86 (1H, d, $J=8.6$ Hz), 7.50 (1H, d, $J=2.8$ Hz), 7.59 (1H, dd, $J=2.8, 8.6$ Hz).

2) 2-Hydroxymethyl-6-nitro-1,4-benzoxazine (1.00 g, 4.76 mmol) obtained in 1) and triethylamine (708 mg, 7.00 mmol) was dissolved in DMF (30 ml). Methanesulfonyl chloride (545 mg, 4.76 mmol) was added to the solution, which was stirred for 30 minutes. 50% Aqueous dimethylamine solution (3 ml) was added to the reaction mixture, which was stirred at 70°C for 4 hours. Ethyl acetate and water were added to the mixture, and extraction was conducted. The organic layer was washed, and concentrated. The residue was subjected to alumina column chromatography, and eluted with ethyl acetate: n-hexane (40:60), to give 2-[(dimethylamino)methyl]-6-nitro-1,4-benzoxazine (790 mg) as a colorless oily substance.

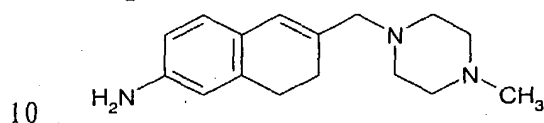
$^1\text{H-NMR}$ (CDCl_3) δ : 2.33 (6H, s), 2.47-2.67 (2H, m), 3.19-3.25 (1H, m), 3.46-3.52 (1H, m), 4.09 (1H, brs), 4.30-4.35 (1H, m), 6.86 (1H, d, $J=8.9$ Hz), 7.48 (1H, d, $J=2.8$ Hz), 7.57 (1H, dd, $J=2.8, 8.9$ Hz).

3) 2-[(Dimethylamino)methyl]-6-nitro-1,4-benzoxazine (760 mg, 3.2 mmol) obtained in 2) was dissolved in methanol (10 ml). Concentrated hydrochloric acid (3 ml) and iron powder (0.80 g) were added to the solution, which was stirred for 2 hours. The reaction mixture was concentrated. 1N Aqueous sodium hydroxide solution and ethyl acetate was added to the residue, and extraction was conducted. The organic layer was concentrated. The residue was subjected to alumina column chromatography, and eluted with ethyl acetate: n-hexane (20:80), to give the

titled compound (430 mg) as a colorless oily substance.
¹H-NMR (CDCl₃) δ: 2.31 (6H, s), 2.41-2.62 (2H, m), 3.12-3.17
(1H, m), 3.36-3.41 (1H, m), 3.30-3.50 (2H, brs), 3.67 (1H,
brs), 4.12-4.21 (1H, m), 5.99 (1H, d, J=2.5 Hz), 6.03 (1H,
5 dd, J=2.5, 8.4 Hz), 6.65 (1H, d, J=8.4 Hz).

Reference Example 106

6-[(4-Methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenamine

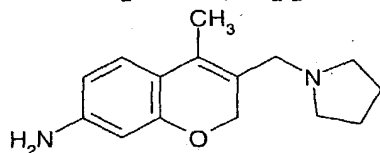


The titled compound was obtained by carrying out the same operation as in Reference Example 52, using 6-acetamido-2-(N,N-dimethylaminomethylidene)-1-tetralone obtained in Example 41-1).

15 ¹H NMR (CDCl₃) δ: 2.27 (2H, t, J=8.1 Hz), 2.29 (3H, s), 2.45 (8H, bs), 2.72 (2H, t, J=8.1 Hz), 3.03 (2H, s), 3.60 (2H, s), 6.26 (1H, s), 6.45-6.47 (2H, m), 6.80-6.83 (1H, m).

Reference Example 107

20 4-Methyl-3-(1-pyrrolidinylmethyl)-2H-chromen-7-amine



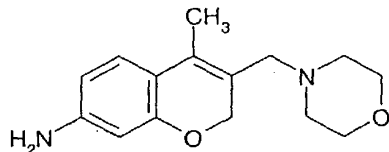
The titled compound was obtained by carrying out the same operations as in Example 41-1) and Reference Example 69 in this order, using 1-acetylamino-3,4-dihydrochromen-1-one.

25 ¹H NMR (CDCl₃) δ: 1.73-1.83 (4H, m), 1.99 (3H, s), 2.46-2.51 (4H, m), 3.22 (2H, s), 3.70 (2H, bs), 4.66 (2H, s), 6.18 (1H, d, J=2.2 Hz), 6.26 (1H, dd, J=2.2 Hz, 8.1 Hz), 7.00 (1H, d, J=8.1 Hz).

30

Reference Example 108

4-Methyl-3-(4-morpholinylmethyl)-2H-chromen-7-amine

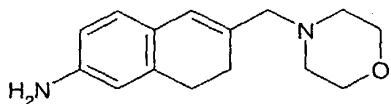


The titled compound was obtained by carrying out the same operations as in Example 41-1) and Reference Example 69 in this order, using 1-acetylamino-3,4-

¹H NMR (CDCl₃) δ: 1.98 (3H, s), 2.41-2.44 (4H, m), 3.08 (2H, s), 3.66-3.69 (6H, m), 4.62 (2H, s), 6.18 (1H, d, J=2.2 Hz), 6.26 (1H, dd, J=2.2 Hz, 8.1 Hz), 7.00 (1H, d, J=8.1 Hz).

Reference Example 109

6-(4-Morpholinylmethyl)-7,8-dihydro-2-naphthalenamine

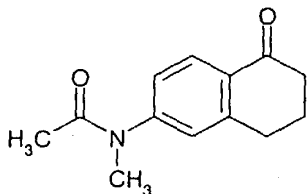


The titled compound was obtained by carrying out the same operations as in Reference Example 52, using 6-acetamido-2-(N,N-dimethylaminomethylidene)-1-tetralone obtained in Example 41-1).

¹H-NMR (CDCl₃) δ: 2.28 (2H, t, J=7.8 Hz), 2.42 (4H, t, J=4.4 Hz), 2.72 (2H, t, J=7.8 Hz), 3.01 (2H, s), 3.60 (2H, brs.), 3.70 (4H, t, J=4.4 Hz), 6.26 (1H, s), 6.46 (2H, m), 6.82 (1H, d, J=8.7 Hz).

Reference Example 110

N-Methyl-N-(5-oxo-5,6,7,8-tetrahydro-2-naphthalenyl)acetamide



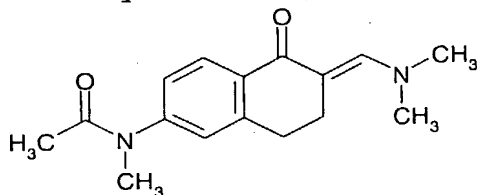
6-Acetamido-1-tetralone (13.7 g, 67.4 mmol) was dissolved in tetrahydrofuran (40 ml). Sodium

hydride(oil)(2.40 g, 101 mmol) was added to the solution, which was refluxed with heating for 2.5 hours. After cooling, methyl iodide(20 ml) was added to the reaction mixture, which was stirred at 40°C for 15 hours. The reaction mixture was poured into a cold water, and extraction was conducted using ethyl acetate. The extract was washed with 1N hydrochloric acid and 1 N aqueous sodium hydroxide solution. The ethyl acetate layer was concentrated. The residue was purified by alumina column chromatography (development solvent; ethyl acetate:n-hexane = 50:50 ~ 100:0) . The eluent was concentrated under reduced pressure. The resulting residue was recrystallized from ethyl acetate - diisopropyl ether, to give the titled compound(8.3 g).

¹H-NMR (CDCl₃) δ: 1.96 (3H, s), 2.19(2H, m), 2.69 (2H, t, J=6.2 Hz), 2.99 (2H, t, J=5.9 Hz), 3.29 (3H, s), 7.10-7.15 (2H, m), 8.09 (1H, d, J=8.4 Hz).

Reference Example 111

N-[6-[(E)-(Dimethylamino)methylidene]-5-oxo-5,6,7,8-tetrahydro-2-naphthalenyl]-N-methylacetamide



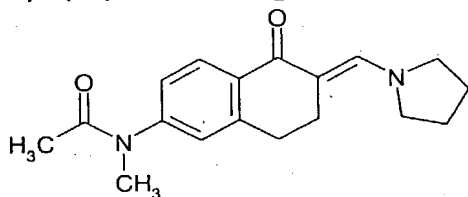
N-Methyl-N-(5-oxo-5,6,7,8-tetrahydro-2-naphthalenyl)acetamide (4.3 g, 19.8 mmol) obtained in Reference Example 110 was dissolved in N,N-dimethylformamide-dimethylacetal(50 ml), which was refluxed with heating under nitrogen atmosphere for 15 hours. The reaction mixture was concentrated under reduced pressure. The resulting residue was washed with ethyl acetate - diisopropyl ether, to give the titled compound(3.9 g).

¹H-NMR (CDCl₃) δ: 1.93 (3H, s), 2.86 (2H, t, J=7.3 Hz), 2.95

(2H, t, $J=7.3$ Hz), 3.16 (6H, s), 3.28 (3H, s), 6.99 (1H, s), 7.09 (1H, d, $J=8.1$ Hz), 7.75 (1H, s), 8.07 (1H, d, $J=8.1$ Hz).

5 Reference Example 112

N-Methyl-N-[5-oxo-6-((E)-1-pyrrolidinylmethylidene)-5,6,7,8-tetrahydro-2-naphthalenyl]acetamide

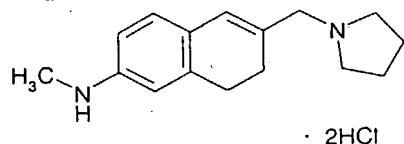


10 N-[6-[(E)-(Dimethylamino)methylidene]-5-oxo-5,6,7,8-tetrahydro-2-naphthalenyl]-N-methylacetamide (5.7 g, 20.9 mmol) obtained in Reference Example 111 was dissolved in pyrrolidine (50 ml), which was refluxed with heating under nitrogen atmosphere for 3.5 hours. The reaction mixture was poured into cold water, and extraction
15 was conducted using ethyl acetate. The ethyl acetate layer was concentrated. The resulting residue was recrystallized from ethyl acetate - diisopropyl ether, to give the titled compound (4.0 g).

¹H-NMR (CDCl₃) δ : 1.93-1.96 (7H, m), 2.85 (2H, t, $J=6.7$ Hz),
20 2.96 (2H, t, $J=6.7$ Hz), 3.28 (3H, s), 3.63 (4H, m), 6.99 (1H, s), 7.10 (1H, dd, $J=8.4$, 2.0 Hz), 7.95 (1H, s), 8.08 (1H, d, $J=8.4$ Hz).

Reference Example 113

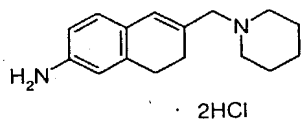
25 N-Methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine dihydrochloride



N-Methyl-N-[5-oxo-6-((E)-1-pyrrolidinylmethylidene)-5,6,7,8-tetrahydro-2-naphthalenyl]acetamide (4.0 g, 13.4 mmol) obtained in
30

Reference Example 112 was dissolved in methanol - acetic acid(10:1, 220 ml). 10% Palladium on carbon (0.4 g) was added to the solution, which was stirred under hydrogen atmosphere for 48 hours. The catalyst was filtered off, and
5 the reaction mixture was concentrated under reduced pressure. Ethyl acetate and 1N hydrochloric acid were added to the residue, and extraction was conducted. After the water layer was made alkaline with 4N aqueous sodium hydroxide solution, extraction was conducted using ethyl
10 acetate. The ethyl acetate layer was concentrated. Tetrahydrofuran - 5N hydrochloric acid (50:50, 200 ml) was added to the resulting residue, which was refluxed with heating for 13 hours. The reaction mixture was concentrated. Ethyl acetate and saturated aqueous sodium
15 carbonate solution was added to the residue, and extraction was conducted. 4N Hydrogen chloride - ethyl acetate solution was added to the ethyl acetate layer, which was concentrated under reduced pressure. The resulting residue was recrystallized from methanol - ethyl acetate,
20 to give the titled compound(2.8 g).
¹H-NMR (DMSO-d₆) δ : 1.98 (4H, m), 2.45 (4H, m), 2.81 (5H, m), 3.01 (2H, m), 3.44 (2H, m), 3.85 (1H, s), 3.86 (1H, s), 6.67 (1H, s), 7.02-7.10 (3H, m), 10.90 (1H, brs.).

25 Reference Example 114
6-(1-Piperidinylmethyl)-7,8-dihydro-2-naphthalenamine dihydrochloride



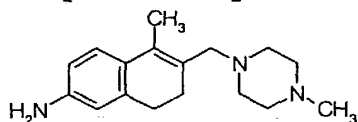
The titled compound was obtained by carrying out the
30 same operation as in Reference Example 52, using 6-acetamido-2-(N,N-dimethylaminomethylidene)-1-tetralone obtained in Example 41-1).

¹H-NMR (DMSO-d₆) δ : 1.39 (1H, m), 1.80 (5H, m), 2.50 (5H, m), 2.83 (4H, m), 3.35-3.38 (2H, m), 3.79 (2H, s), 6.70 (1H,

s), 7.05-7.13 (3H, m), 10.40 (1H, brs).

Reference Example 115

5-Methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenamine

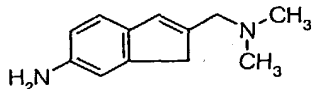


The titled compound was obtained by carrying out the same operation as in Reference Example 69, using 6-acetamido-2-(N,N-dimethylaminomethylidene)-1-tetralone obtained in Example 41-1).

¹H NMR (CDCl₃) δ: 2.02 (3H, s), 2.27 (2H, t, J=8.1 Hz), 2.27 (3H, s), 2.44 (8H, bs), 2.63 (2H, t, J=8.1 Hz), 3.12 (2H, s), 3.61 (2H, s), 6.48-6.54 (2H, m), 7.08 (1H, d, J=7.8 Hz).

15. Reference Example 116

2-[(Dimethylamino)methyl]-1H-inden-6-amine



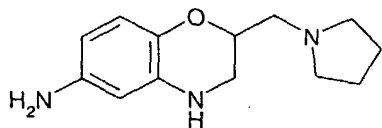
The titled compound was obtained by carrying out the same operation as in Example 41-2), using N-[2-[(E)-(dimethylamino)methylidene]-1-oxo-2,3-dihydro-1H-inden-5-yl]acetamide obtained in Reference Example 47.

¹H NMR (CDCl₃) δ: 2.24 (6H, s), 3.26 (2H, s), 3.33 (2H, s), ca.3.5 (2H, br), 6.58 (2H, m), 6.81 (1H, s), 7.08 (1H, d, J=8.1 Hz).

25

Reference Example 117

6-Amino-2-(1-pyrrolidinylmethyl)-3,4-dihydro-2H-1,4-benzoxazine



30

A mixture of 6-nitro-2-(1-pyrrolidinylmethyl)-3,4-dihydro-2H-1,4-benzoxazine and 4-(methylsulfonyl)-

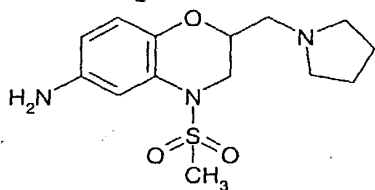
6-nitro-2-(1-pyrrolidinylmethyl)-3,4-dihydro-2H-1,4-benzoxazine was obtained by carrying out the same operation as in Reference Example 105-2), using 2-hydroxymethyl-6-nitro-3,4-dihydro-2H-1,4-benzoxazine obtained in
 5 Reference Example 105-1).

The titled compound was obtained by carrying out the same operation as in Reference Example 105-3), using the mixture obtained above.

¹H-NMR (CDCl₃) δ: 1.76-1.81 (4H, m), 2.50-2.70 (4H, m), 2.70
 10 (2H, d, J=6.3Hz), 3.13-3.20 (1H, m), 3.20-3.40 (2H, brs), 3.39-3.43 (1H, m), 3.66 (1H, brs), 4.11-4.21 (1H, m), 5.99 (1H, d, J=2.7Hz), 6.03 (1H, dd, J=2.7, 8.4 Hz), 6.64 (1H, d, J=8.4 Hz).

15 Reference Example 118

6-Amino-4-(methylsulfonyl)-2-(1-pyrrolidinylmethyl)-3,4-dihydro-2H-1,4-benzoxazine

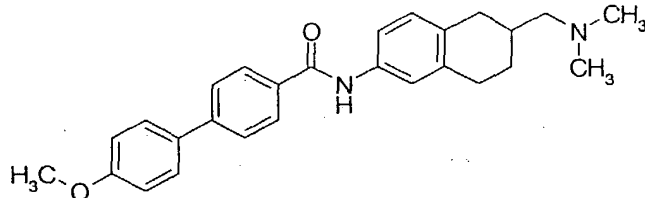


The titled compound was obtained by carrying out the
 20 same operation as in Reference Example 105-3), using the mixture of 6-nitro-2-(1-pyrrolidinylmethyl)-3,4-dihydro-2H-1,4-benzoxazine and 4-(methylsulfonyl)-6-nitro-2-(1-pyrrolidinylmethyl)-3,4-dihydro-2H-1,4-benzoxazine obtained in Reference Example 117.

25 ¹H-NMR (CDCl₃) δ: 1.70-1.80 (4H, m), 2.50-2.70 (4H, m), 2.73 (2H, d, J=6.0Hz), 2.95 (3H, s), 3.21-3.29 (1H, m), 2.80-3.10 (2H, brs), 4.10-4.21 (1H, m), 4.26-4.32 (1H, m), 6.43 (1H, dd, J=2.7, 8.4 Hz), 6.77 (1H, d, J=8.4 Hz), 7.11 (1H, d, J=2.7Hz).

Example 1

N-[2-(N,N-Dimethylamino)methyl-6-tetralinyl]-(4'-methoxybiphenyl-4-yl)carboxamide



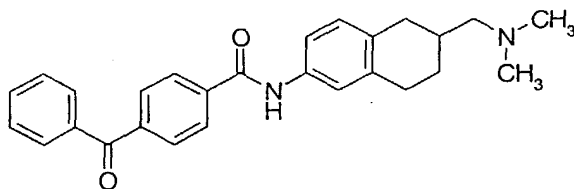
5 DMF solution (0.25 ml) of 2M HOBt, DMF solution (0.30 ml) of 2M WSCD, triethylamine (0.14 ml) and DMAP (0.132 g) were added to DMF solution (3 ml) of 6-amino-2-(N,N-dimethylamino)methyltetralin (0.139 g) and 4-(4-methoxyphenyl)benzoic acid (0.118 g). After the reaction mixture
10 was stirred at room temperature for 12 hours, 10% potassium carbonate solution was added, and extraction was conducted using ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The resulting crude crystal
15 was washed with diethyl ether, which was recrystallized using ethyl acetate-hexane, to give the titled compound (0.124 g).

Melting point: 170 - 175°C.

20 Compounds described in the following Examples 2 and 3 were produced in the same manner as in Example 1.

Example 2

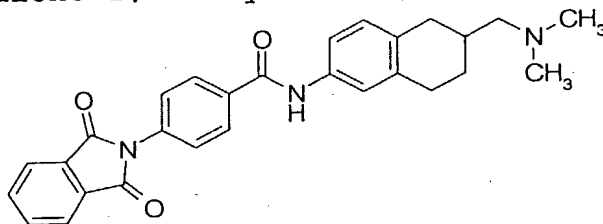
25 4-Benzoyl-N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]benzamide



Melting point: 193 - 196°C (recrystallization solvent: ethyl acetate-hexane)

Example 3

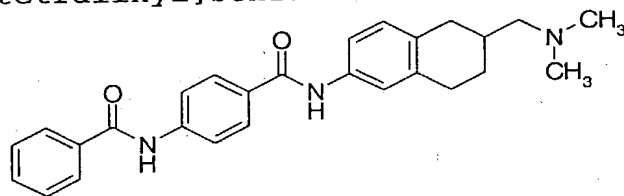
N-[2-(N,N-Dimethylamino)methyl-6-tetralinyl]-4-(1,3-dioxo-1,3-dihydro-2H-isoindol-2-yl) benzamide



5 Melting point: 235 - 240°C (washed with diethyl ether)

Example 4

4-(Benzoylamino)-N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]benzamide



10

6-Amino-2-(N,N-dimethylamino)methyltetralin hydrochloride (139 mg), 4-benzoylamino benzoic acid (121 mg), WSCD (0.13 ml), HOBt (92 mg), triethylamine (0.14 ml) and DMAP (61 mg) were added to DMF (4 ml). After the reaction mixture was shaken at room temperature for 20 hours using a shaker, the reaction mixture was poured into water, and extraction was conducted using ethyl acetate-THF (1:1).

15 The organic layer was washed with water, saturated sodium bicarbonate solution and saturated aqueous sodium chloride solution, dried, and then concentrated. The resulting crude crystal was washed with hexane, to give the titled compound (181 mg).

20 Melting point : 241 - 242°C

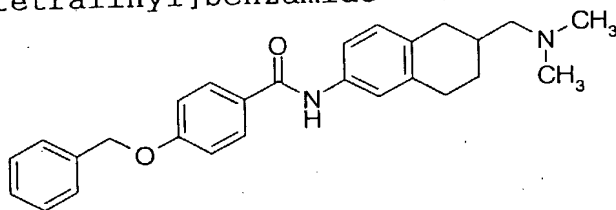
Washing solvent : hexane

25

Compounds described in the following Examples 5 to 14 were produced in the same manner as in Example 4.

Example 5

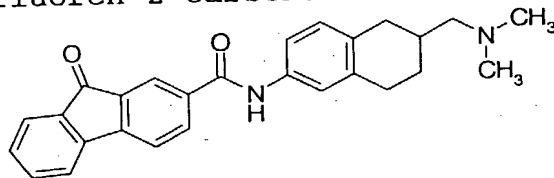
4-(Benzyloxy)-N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]benzamide



- 5 Melting point : 135 - 136°C
Washing solvent : hexane

Example 6

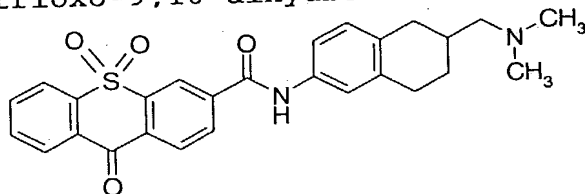
- 10 N-[2-(N,N-Dimethylamino)methyl-6-tetralinyl]-9-oxo-9H-fluoren-2-carboxamide



- Melting point : 224 - 226°C
Washing solvent : hexane

15 Example 7

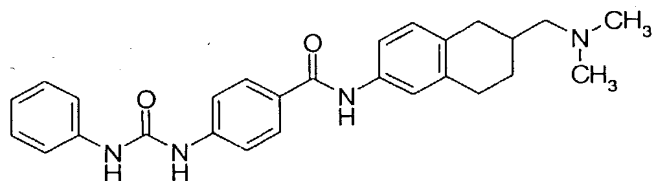
N-[2-(N,N-Dimethylamino)methyl-6-tetralinyl]-9,10,10-trioxo-9,10-dihydro-101⁶-thioxanthene-3-carboxamide



- Melting point : 222 - 223°C (decomposition)
20 Washing solvent: hexane

Example 8

(4-Anilinocarbonyl)amino-N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]benzamide

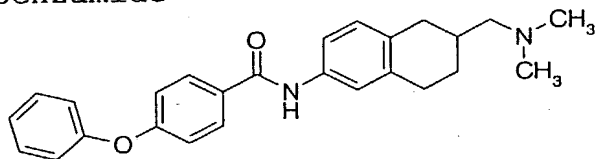


Melting point : 216 - 217°C (decomposition)

Washing solvent : hexane

5 Example 9

N-[2-(N,N-Dimethylamino)methyl-6-tetralinyl]-4-phenoxy
benzamide

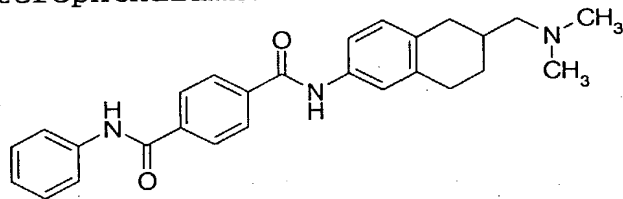


Melting point : 137 - 139°C

10 Washing solvent : hexane

Example 10

N¹-[2-(N,N-Dimethylamino)methyl-6-tetralinyl]-N⁴-phenyl
terephthalamide



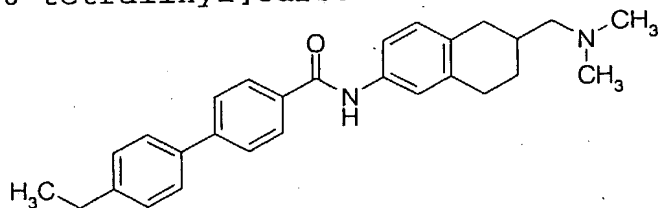
15

Melting point : 238 - 240°C (decomposition)

Washing solvent : hexane

Example 11

20 (4'-Ethylbiphenyl-4-yl)-N-[2-(N,N-dimethylamino)methyl-
6-tetralinyl]carboxamide

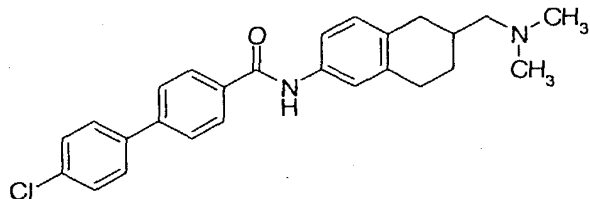


Melting point : 137 - 138°C

Washing solvent : hexane

Example 12

(4'-Chlorobiphenyl-4-yl)-N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]carboxamide

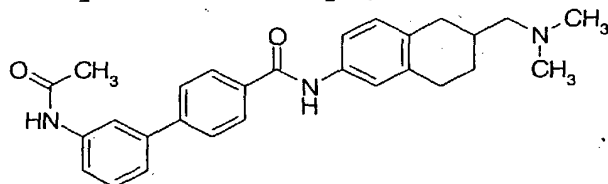


Melting point : 187 - 189°C

Washing solvent : hexane

Example 13

(4'-Acetylaminobiphenyl-4-yl)-N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]carboxamide

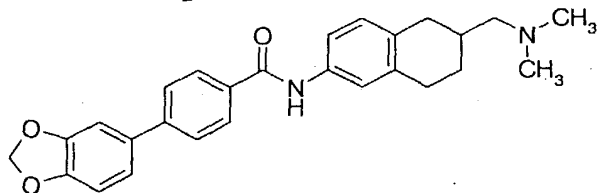


Melting point : 183 - 186°C

Washing solvent : hexane

Example 14

4-(1,3-Benzodioxol-5-yl)-N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]benzamide



Melting point : 174 - 176°C

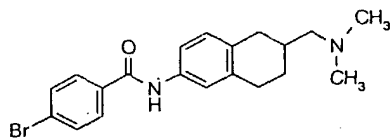
Washing solvent : hexane

Example 15

4-Bromo-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-

tetrahydro-2-naphthalenyl]benzamide

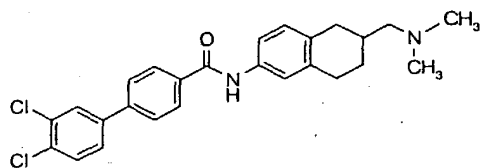
The titled compound was obtained as a white powder by the same method as in Example 1.



5 Melting point: 141 - 143°C (washing solvent: n-hexane)

Example 16

3',4'-Dichloro-N-[6-[(N,N-dimethylamino)methyl]-
5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-
10 carboxamide



4-Bromo-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-
tetrahydro-2-naphthalenyl]benzamide (400 mg, 1.03 mmol)
obtained in Example 15, 3,4-dichlorophenylboric acid (50
15 wt% THF-H₂O solution, 0.473 ml, 1.24 mmol), and 2N sodium
carbonate solution (1.03 ml, 2.07 mmol) were dissolved in
50 ml of dimethoxyethane, then palladium
tetrakis(triphenylphosphine) (35.8 mg, 0.031 mmol) was added
under nitrogen atmosphere, which was stirred at 90°C for
20 15 hours.

Ethyl acetate was added to the reaction mixture, which
was washed with saturated aqueous sodium chloride solution,
dried using anhydrous magnesium sulfate, and the solvent
was distilled out under reduced pressure. The residue was
25 refined by alumina column chromatography (development
solvent; n-hexane:ethyl acetate = 3:1), and pulverized with
n-hexane to give the titled compound (204 mg) a white
powder.

¹H-NMR (CDCl₃) δ: 1.41 (1H, m), 1.95 (2H, m), 2.26 (6H, s),
30 2.26-2.45 (3H, m), 2.83-2.99 (3H, m), 7.10 (1H, d, J=8.1
Hz), 7.26-7.77 (8H, m), 7.94 (2H, d, J=8.4 Hz).

Elemental analysis for $C_{26}H_{26}Cl_2N_2O \cdot 0.1H_2O$

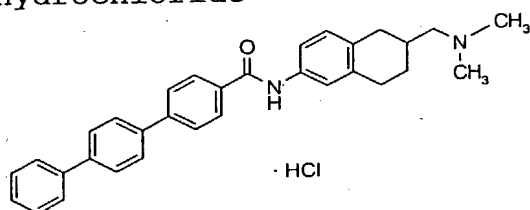
Calcd.: C, 68.60; H, 5.80; N, 6.15.

Found: C, 68.42; H, 5.60; N, 5.92.

Melting point: 143 - 145°C (crystallization solvent:
ethyl acetate-hexane)

Example 17

N-[6-[(N,N-Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-4'-phenyl[1,1'-biphenyl]-4-carboxamide
hydrochloride



The free basic substance (35 mg) of the titled compound was obtained in the same manner as in Example 16, using 4-bromo-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]benzamide (400 mg, 1.03 mmol) obtained in Example 15, and 4-biphenylboric acid (1.25 g, 1.25 mmol). The resulting free basic substance (30 mg) was dissolved in 10 ml of methanol, then 100 ml of 1N hydrochloric acid was added, and the reaction mixture was stirred. The reaction mixture was concentrated, and pulverized using diethyl ether, to give the titled compound (35.3 mg) as a white powder.

1H -NMR (DMSO- d_6 , free base) δ : 1.32 (1H, m), 1.93 (2H, m), 2.15 (6H, s), 2.15-2.36 (3H, m), 2.74-2.94 (3H, m), 7.05 (1H, d, $J=8.4$ Hz), 7.40-7.55 (5H, m), 7.73-7.91 (8H, m), 8.07 (2H, d, $J=8.4$ Hz), 10.14 (1H, s).

Elemental analysis for $C_{32}H_{32}N_2O \cdot HCl \cdot 2H_2O$

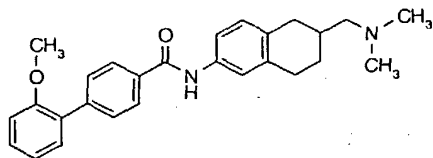
Calcd.: C, 72.10; H, 7.00; N, 5.25.

Found: C, 71.81; H, 6.57; N, 5.08.

Melting point: 220°C (decomposition) (crystallization solvent: methanol-diethyl ether)

Example 18

N-[6-[(N,N-Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-2'-methoxy[1,1'-biphenyl]-4-carboxamide



5 The titled compound (208 mg) was obtained as a white powder by the same method as in Example 16, using 4-bromo-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]benzamide (250 mg, 0.645 mmol) obtained in Example 15, and 2-methoxyphenylboric acid (118
10 mg, 0.775 mmol).

$^1\text{H-NMR}$ (CDCl_3) δ : 1.42 (1H, m), 1.96 (2H, m), 2.23 (6H, s), 2.23-2.47 (3H, m), 2.85 (3H, m), 3.83 (3H, s), 7.05 (3H, m), 7.34 (3H, m), 7.47 (1H, s), 7.64 (2H, d, $J=8.4$ Hz), 7.79 (1H, s), 7.90 (2H, d, $J=8.4$ Hz).

15 Elemental analysis for $\text{C}_{27}\text{H}_{30}\text{N}_2\text{O}_2 \cdot 0.1\text{H}_2\text{O}$

Calcd.: C, 77.89; H, 7.31; N, 6.73.

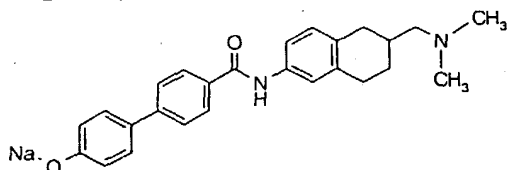
Found: C, 77.86; H, 7.18; N, 6.79.

Melting point: 155 - 157°C (crystallization solvent: ethyl acetate-hexane)

20

Example 19

Sodium salt of N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-4'-oxy[1,1'-biphenyl]-4-carboxamide



25

The titled compound (117 mg) was obtained as a white powder by the same method as in Example 16, using 4-bromo-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]benzamide (250 mg, 0.645 mmol) and 4-hydroxyphenylboric acid (107 mg, 0.775 mmol).
30

$^1\text{H-NMR}$ (DMSO-d_6) δ : 1.36 (1H, m), 1.89 (2H, m), 2.15 (6H, s), 2.15-2.35 (3H, m), 2.77 (3H, m), 6.88 (2H, d, $J=8.4$ Hz), 7.02 (1H, d, $J=8.4$ Hz), 7.48 (1H, d, $J=8.4$ Hz), 7.53 (1H, s), 7.59 (2H, d, $J=8.4$ Hz), 7.73 (2H, d, $J=8.4$ Hz), 8.00 (2H, d, $J=8.4$ Hz), 10.07 (1H, s).

Elemental analysis for $\text{C}_{26}\text{H}_{27}\text{N}_2\text{O}_2\text{Na} \cdot 0.2\text{H}_2\text{O}$

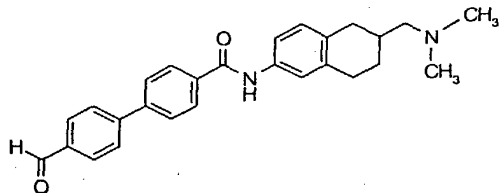
Calcd.: C, 73.29; H, 6.48; N, 6.59.

Found: C, 73.25; H, 6.18; N, 6.36.

Melting point: 246 - 248°C (crystallization solvent: ethyl acetate-diethyl ether)

Example 20

N-[6-[(N,N-Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-4'-formyl[1,1'-biphenyl]-4-carboxamide



15

The titled compound (205 mg) was obtained as a white powder by the same method as in Example 16, using 4-bromo-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]benzamide (250 mg, 0.645 mmol) and 4-formylphenylboric acid (145 mg, 0.968 mmol).

$^1\text{H-NMR}$ (CDCl_3) δ : 1.41 (1H, m), 1.95 (2H, m), 2.26 (6H, s), 2.26-2.42 (3H, m), 2.85-2.94 (3H, m), 7.09 (2H, d, $J=8.1$ Hz), 7.32 (1H, d, $J=8.4$ Hz), 7.47 (1H, m), 7.63-7.94 (3H, m), 7.87-7.99 (4H, m), 8.13 (1H, s), 10.11 (1H, s).

Elemental analysis for $\text{C}_{27}\text{H}_{28}\text{N}_2\text{O}_2 \cdot 0.2\text{H}_2\text{O}$

Calcd.: C, 77.93; H, 6.88; N, 6.73.

Found: C, 77.89; H, 6.75; N, 6.71.

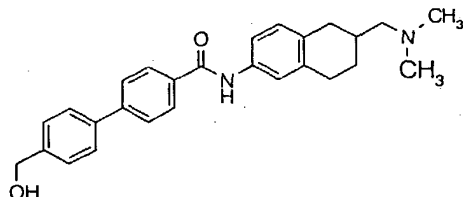
Melting point: 130 - 132°C (crystallization solvent: ethyl acetate-diethyl ether)

30

Example 21

N-[6-[(N,N-Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-

naphthalenyl]-4'-(hydroxymethyl)[1,1'-biphenyl]-4-carboxamide



5 N-[6-[(N,N-Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-4'-formyl[1,1'-biphenyl]-4-carboxamide (100 mg, 0.242 mmol) was dissolved in tetrahydrofuran-methanol (1:1) solution (2.4 ml), then sodium borohydride (18.3 mg, 0.485 mmol) was added, which was stirred for 2 hours. Ethyl acetate was added to the
10 reaction mixture, which was washed with saturated aqueous sodium chloride solution, dried using anhydrous magnesium sulfate, and the solvent was distilled out under reduced pressure. The residue was pulverized using ether-n-hexane, to give the titled compound (86 mg) as a white
15 powder.

¹H-NMR (CDCl₃) δ: 1.39 (1H, m), 1.94 (2H, m), 2.25 (6H, s), 2.25-2.44 (3H, m), 2.82-2.95 (3H, m), 4.78 (2H, s), 7.07 (1H, d, J=8.4 Hz), 7.31 (1H, d, J=8.4 Hz), 7.38-7.56 (4H, m), 7.64-7.70 (3H, m), 7.85 (1H, s), 7.93 (2H, d, J=8.4 Hz).

20 Elemental analysis for C₂₇H₃₀N₂O₂ · 0.2H₂O

Calcd.: C, 77.56; H, 7.33; N, 6.70.

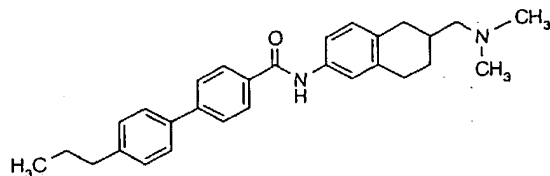
Found: C, 77.53; H, 7.27; N, 6.55.

Melting point: 138 - 139°C (crystallization solvent: ethyl acetate-diethyl ether)

25

Example 22

N-[6-[(N,N-Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-4'-propyl[1,1'-biphenyl]-4-carboxamide



The titled compound (158 mg) was obtained as a white powder by the same method as in Example 1, using N-[(6-amino-1,2,3,4-tetrahydro-2-naphthalenyl)methyl]-N,N-dimethylamine (102 mg, 0.499 mmol), and 4-(4-propyl)benzoic acid (144 mg, 0.599 mmol).

¹H-NMR (CDCl₃) δ: 0.98 (3H, t, J=7.5 Hz), 1.40 (1H, m), 1.69 (2H, m), 1.94 (2H, m), 2.25 (6H, s), 2.25-2.45 (3H, m), 2.64 (2H, t, J=7.5 Hz), 2.85 (3H, m), 7.08 (1H, d, J=7.8 Hz), 7.26 (3H, m), 7.46 (1H, s), 7.54 (2H, d, J=8.1 Hz), 7.67 (2H, d, J=8.1 Hz), 7.81 (1H, s), 7.91 (2H, d, J=8.4 Hz).
Elemental analysis for C₂₉H₃₄N₂O

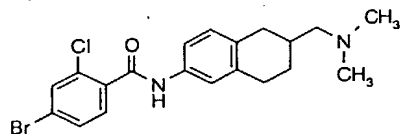
Calcd.: C, 81.65; H, 8.03; N, 6.57.

Found: C, 81.30; H, 7.94; N, 6.40.

Melting point: 186 - 188°C (crystallization solvent: ethyl acetate-diethyl ether)

Example 23

4-Bromo-2-chloro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]benzamide



The titled compound (483 mg) was obtained as a white powder by the same method as in Example 1, using N-[(6-amino-1,2,3,4-tetrahydro-2-naphthalenyl)methyl]-N,N-dimethylamine (300 mg, 1.47 mmol) and 4-bromo-2-chloro benzoic acid (415 mg, 1.76 mmol).

¹H-NMR (CDCl₃) δ: 1.40 (1H, m), 1.94 (2H, m), 2.25 (6H, s), 2.25-2.44 (3H, m), 2.94 (3H, m), 7.08 (1H, d, J=8.4 Hz), 7.28 (1H, m), 7.41 (1H, s), 7.50 (1H, m), 7.61 (2H, m), 7.81 (1H, s).

Elemental analysis for C₂₀H₂₂BrClN₂O

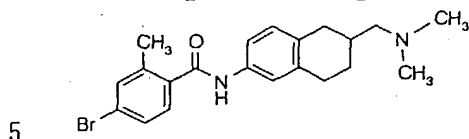
Calcd.: C, 56.96; H, 5.26; N, 6.64.

Found: C, 57.09; H, 5.37; N, 6.55.

Melting point: 130 - 132°C (crystallization solvent: ethyl acetate-diethyl ether)

Example 24

4-Bromo-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-2-methylbenzamide



The titled compound (418 mg) was obtained as a white powder by the same method as in Example 1, using N-[(6-amino-1,2,3,4-tetrahydro-2-naphthalenyl)methyl]-N,N-dimethylamine (293 mg, 1.43 mmol) and 4-bromo-2-methyl

10 benzoic acid (370 mg, 1.72 mmol).

$^1\text{H-NMR}$ (CDCl_3) δ : 1.40 (1H, m), 2.04 (2H, m), 2.25 (6H, s), 2.25-2.40 (3H, m), 2.46 (3H, s), 2.88 (3H, m), 7.07 (1H, d, $J=7.8$ Hz), 7.21-7.41 (6H, m).

Elemental analysis for $\text{C}_{21}\text{H}_{25}\text{BrN}_2\text{O}$

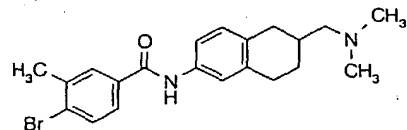
15 Calcd.: C, 62.85; H, 6.28; N, 6.98.

Found: C, 63.10; H, 6.11; N, 6.97.

Melting point: 140 - 142°C (crystallization solvent: ethyl acetate-hexane)

20 Example 25

4-Bromo-N-[6[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-3-methylbenzamide



The titled compound (434 mg) was obtained as a white powder by the same method as in Example 1, using N-[(6-amino-1,2,3,4-tetrahydro-2-naphthalenyl)methyl]-N,N-dimethylamine (300 mg, 1.47 mmol) and 4-bromo-3-methyl benzoic acid (379 mg, 1.76 mmol).

25

$^1\text{H-NMR}$ (CDCl_3) δ : 1.40 (1H, m), 1.93 (2H, m), 2.25 (6H, s), 2.25-2.40 (3H, m), 2.46 (3H, s), 2.87 (3H, m), 7.07 (1H, d, $J=7.8$ Hz), 7.29 (1H, m), 7.40 (1H, s), 7.49 (1H, m), 7.61 (1H, d, $J=8.1$ Hz), 7.72 (2H, s-like).

30

Elemental analysis for $C_{21}H_{25}BrN_2O$

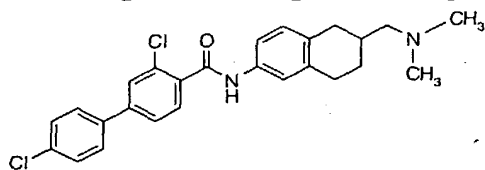
Calcd.: C, 62.85; H, 6.28; N, 6.98.

Found: C, 62.84; H, 6.05; N, 6.93.

Melting point: 154 - 155°C (crystallization solvent: ethyl acetate-hexane)

Example 26

3,4'-Dichloro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



The titled compound (122 mg) was obtained as a white powder by the same method as in Example 16, using 4-bromo-2-chloro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]benzamide (250 mg, 0.607 mmol) obtained in Example 23, and 4-chlorophenyl boric acid (114 mg, 0.729 mmol).

1H -NMR ($CDCl_3$) δ : 1.41 (1H, m), 1.95 (2H, m), 2.26 (6H, s), 2.26-2.42 (3H, m), 2.85 (3H, m), 7.10 (1H, d, $J=8.4$ Hz), 7.31 (1H, m), 7.43-7.63 (8H, m), 7.87 (1H, d, $J=8.1$ Hz).

Elemental analysis for $C_{26}H_{26}Cl_2N_2O$

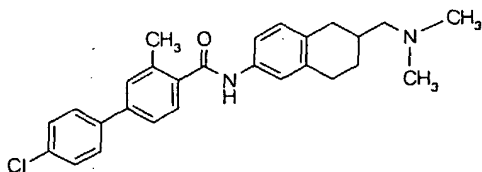
Calcd.: C, 68.87; H, 5.78; N, 6.18.

Found: C, 68.61; H, 5.49; N, 6.10.

Melting point: 177 - 179°C (crystallization solvent: ethyl acetate-diethyl ether)

Example 27

4'-Chloro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-3-methyl[1,1'-biphenyl]-4-carboxamide



The titled compound (129 mg) was obtained as a white powder by the same method as in Example 16, using 4-bromo-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-2-methylbenzamide (250 mg, 0.623 mmol) obtained in Example 24, and 4-chlorophenylboric acid (117 mg, 0.747 mmol).

$^1\text{H-NMR}$ (CDCl_3) δ : 1.42 (1H, m), 1.96 (2H, m), 2.37 (6H, s), 2.37-2.47 (3H, m), 2.56 (3H, s), 2.90 (3H, m), 7.08 (1H, d, $J=8.1$ Hz), 7.26 (1H, m), 7.41 (6H, m), 7.53 (3H, m).

Elemental analysis for $\text{C}_{27}\text{H}_{29}\text{ClN}_2\text{O} \cdot \text{H}_2\text{O}$

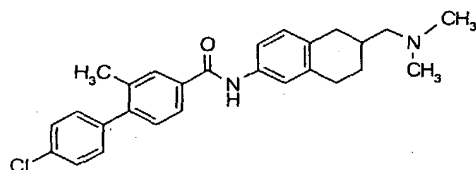
Calcd.: C, 71.90; H, 6.93; N, 6.21.

Found: C, 71.92; H, 6.52; N, 5.92.

Melting point: 163 - 165°C (crystallization solvent: ethyl acetate-diethyl ether)

Example 28

4'-Chloro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-2-methyl[1,1'-biphenyl]-4-carboxamide



The titled compound (168 mg) was obtained as a white powder by the same method as in Example 16, using 4-bromo-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-3-methylbenzamide (250 mg, 0.623 mmol) obtained in Example 25, and 4-chlorophenylboric acid (117 mg, 0.747 mmol).

$^1\text{H-NMR}$ (CDCl_3) δ : 1.41 (1H, m), 1.95 (2H, m), 2.26 (6H, s), 2.24-2.42 (3H, m), 2.33 (3H, s), 2.85 (3H, m), 7.09 (1H, d, $J=8.4$ Hz), 7.26 (4H, m), 7.43 (3H, m), 7.73 (3H, m).

Elemental analysis for $\text{C}_{27}\text{H}_{29}\text{ClN}_2\text{O} \cdot 0.2\text{H}_2\text{O}$

Calcd.: C, 74.28; H, 6.79; N, 6.42.

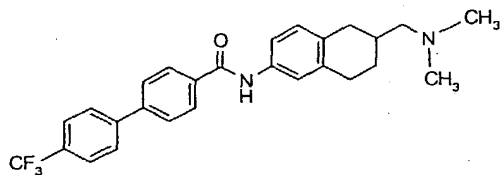
Found: C, 74.27; H, 6.73; N, 6.27.

Melting point: 193 - 195°C (crystallization solvent: ethyl

acetate-diethyl ether)

Example 29

N-[6-[(N,N-Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-4'-(trifluoromethyl)[1,1-biphenyl]-4-carboxamide



The titled compound (194 mg) was obtained as a white powder by the same method as in Example 16, using 4-bromo-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]benzamide (250 mg, 0.645 mmol) obtained in Example 15, and 4-trifluoromethylphenylboric acid (147 mg, 0.775 mmol).

¹H-NMR (CDCl₃) δ: 1.41 (1H, m), 1.95 (2H, m), 2.25 (6H, s), 2.25-2.45 (3H, m), 2.89 (3H, m), 7.09 (1H, d, J=8.1 Hz), 7.31 (1H, d, J=8.1 Hz), 7.46 (1H, s), 7.70 (6H, m), 7.80 (1H, m), 7.96 (2H, d, J=8.4 Hz).

Elemental analysis for C₂₇H₂₇F₃N₂O

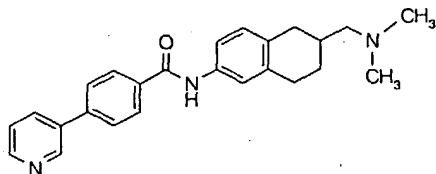
Calcd.: C, 71.66; H, 6.01; N, 6.19.

Found: C, 71.44; H, 6.05; N, 6.09.

Melting point: 205 - 206°C (crystallization solvent: ethyl acetate-diisopropyl ether)

Example 30

N-[6-[(N,N-Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-4-(3-pyridinyl)benzamide



The titled compound (194 mg) was obtained as a white powder by the same method as in Example 16, using 4-bromo-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-

tetrahydro-2-naphthalenyl)benzamide (250 mg, 0.645 mmol) obtained in Example 15, and 2-(3-pyridyl)-1,3,2,-dioxaborinane (126 mg, 0.775 mmol).

¹H-NMR (CDCl₃) δ: 1.41 (1H, m), 1.95 (2H, m), 2.26 (6H, s), 2.26-2.42 (3H, m), 2.85 (3H, m), 7.09 (1H, d, J=7.8 Hz), 7.30-7.47 (3H, m), 7.69 (2H, d, J=8.4 Hz), 7.86-7.99 (4H, m), 8.64 (1H, m), 8.87 (1H, m).

Elemental analysis for C₂₅H₂₇N₃O · 0.1H₂O

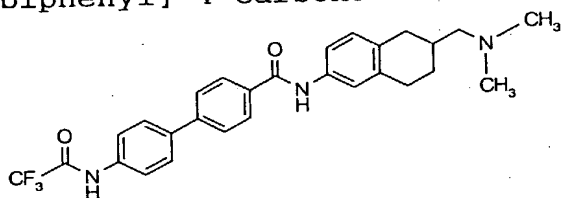
Calcd.: C, 77.53; H, 7.08; N, 10.85.

Found: C, 77.42; H, 7.05; N, 10.58.

Melting point: 177 - 178°C (crystallization solvent: ethyl acetate-diisopropyl ether)

Example 31

N-[6-[(N,N-Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-4'-[(trifluoroacetyl)amino][1,1'-biphenyl]-4-carboxamide



The titled compound (1.02 g) was obtained as a white powder by the same method as in Example 16, using 4-bromo-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]benzamide (1.00 g, 2.58 mmol) obtained in Example 15, and 4-trifluoroacetamidophenylboric acid (722 mg, 3.10 mmol). ¹H-NMR (CDCl₃) δ: 1.41 (1H, m), 2.05 (2H, m), 2.26 (6H, s), 2.26-2.42 (3H, m), 2.89 (3H, m), 7.09 (1H, d, J=8.4 Hz), 7.29 (2H, m), 7.46 (1H, s), 7.69 (7H, m), 7.94 (2H, d, J=8.1 Hz).

Elemental analysis for C₂₈H₂₈F₃N₃O₂

Calcd.: C, 67.87; H, 5.70; N, 8.48.

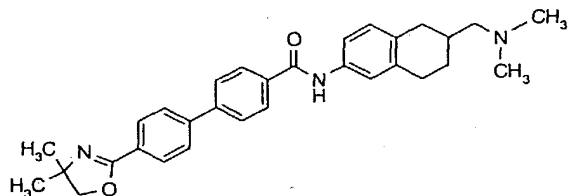
Found: C, 67.70; H, 5.53; N, 8.42.

Melting point: 235 - 237°C (crystallization solvent: ethyl

acetate-diisopropyl ether)

Example 32

N-[6-[(N,N-Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-4'-(4,4-dimethyl-4,5-dihydro-1,3-oxazole-2-yl)[1,1'-biphenyl]-4-carboxamide



The titled compound (238 mg) was obtained as a white powder by the same method as in Example 16, using 4-bromo-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]benzamide (250 mg, 0.645 mmol) obtained in Example 15, and 4-(4,4-dimethyl-4,5-dihydro-1,3-oxazol-2-yl)phenylboronic acid (170 mg, 0.775 mmol).

¹H-NMR (CDCl₃) δ: 1.41 (7H, m), 1.94 (2H, m), 2.25 (6H, s), 2.25-2.41 (3H, m), 2.84 (3H, m), 4.14 (2H, s), 7.08 (1H, d, J=7.8 Hz), 7.30 (1H, m), 7.46 (1H, s), 7.68 (5H, m), 7.94 (2H, d, J=8.4 Hz), 8.03 (2H, d, J=8.4 Hz).

Elemental analysis for C₃₁H₃₅N₃O₂ · 0.2H₂O

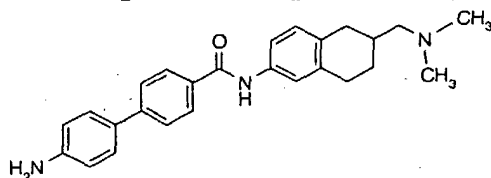
Calcd.: C, 76.74; H, 7.35; N, 8.66.

Found: C, 76.70; H, 7.19; N, 8.49.

Melting point: 185 - 187°C (crystallization solvent: ethyl acetate-diisopropyl ether)

Example 33

4'-Amino-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



N-[6-[(N,N-Dimethylamino)methyl]-5,6,7,8-

tetrahydro-2-naphthalenyl]-4'-
[(trifluoroacetyl)amino][1,1'-biphenyl]-4-carboxamide
(850 mg, 1.72 mmol) obtained in Example 31 was suspended
in a mixed solution of methanol (8 ml) and tetrahydrofuran
5 (4 ml), then 1N sodium hydroxide (3.4 ml) was added, which
was stirred at 50° C for 16 hours. The solvent was distilled
out under reduced pressure, and the residue was pulverized
using water, to give the titled compound (685 mg) as a white
powder.

10 ¹H-NMR (CDCl₃) δ: 1.31 (1H, m), 1.89 (2H, m), 2.15 (6H, s),
2.15-2.34 (3H, m), 2.83 (3H, m), 5.36 (2H, s), 6.67 (2H,
d, J=8.4 Hz), 7.03 (1H, d, J=8.1 Hz), 7.48 (4H, m), 7.68
(2H, d, J=8.1 Hz), 7.96 (2H, d, J=8.4 Hz), 10.02 (1H, s).
Elemental analysis for C₂₆H₂₉N₃O · 1.1H₂O

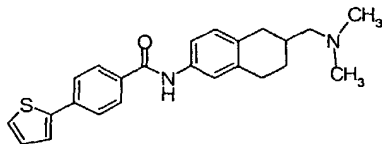
15 Calcd.: C, 74.47; H, 7.50; N, 10.02.

Found: C, 74.39; H, 7.41; N, 9.82.

Melting point: 148 - 150° C (crystallization solvent:
methanol-water)

20 Example 34

N-[6-[(N,N-Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-
naphthalenyl]-4-(2-thienyl) benzamide



The titled compound (70 mg) was obtained as a white
25 powder by the same method as in Example 16, using 4-
bromo-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-
tetrahydro-2-naphthalenyl]benzamide (250 mg, 0.645 mmol)
obtained in Example 15, and 2-thienylboric acid (99.1 mg,
0.775 mmol).

30 ¹H-NMR (CDCl₃) δ: 1.41 (1H, m), 1.94 (2H, m), 2.25 (6H, s),
2.25-2.45 (3H, m), 2.89 (3H, m), 7.11 (2H, m), 7.29-7.45
(4H, m), 7.71 (3H, m), 7.87 (2H, d, J=8.4 Hz).

Elemental analysis for C₂₄H₂₆N₂OS

Calcd.: C, 73.81; H, 6.71; N, 7.17.

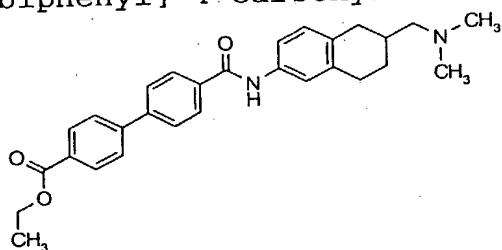
Found: C, 73.49; H, 6.59; N, 7.14.

Melting point: 165 - 166°C (crystallization solvent: ethyl acetate-diisopropyl ether)

5

Example 35

Ethyl 4'-[[[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]amino]carbonyl][1,1'-biphenyl]-4-carboxylate



10

The titled compound (202 mg) was obtained as a white powder by the same method as in Example 16, using 4-bromo-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]benzamide (250 mg, 0.645 mmol) obtained in Example 15, and 4-ethoxycarbonylphenylboric acid (150 mg, 0.775 mmol).

¹H-NMR (CDCl₃) δ: 1.42 (4H, m), 1.95 (2H, m), 2.26 (6H, s), 2.26-2.42 (3H, m), 2.89 (3H, m), 4.41 (2H, q, J=7.2 Hz), 7.09 (1H, d, J=8.4 Hz), 7.31 (1H, d, J=8.4 Hz), 7.47 (1H, s), 7.70 (4H, m), 7.80 (1H, s), 7.96 (2H, d, J=8.4 Hz), 8.14 (2H, d, J=8.4 Hz).

20

Elemental analysis for C₂₉H₃₂N₂O₃

Calcd.: C, 76.29; H, 7.06; N, 6.14.

Found: C, 76.25; H, 7.07; N, 6.09.

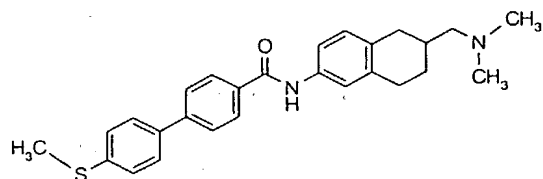
Melting point: 156 - 158°C (crystallization solvent: ethyl acetate-diisopropyl ether)

25

Example 36

N-[6-[(N,N-Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-4'-(methylsulfanyl)[1,1'-biphenyl]-4-carboxamide

30



The titled compound (360 mg) was obtained as a white powder by the same method as in Example 16, using 4-bromo-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]benzamide (500 mg, 1.29 mmol) obtained in Example 15, and 4-methylthiophenylboric acid (260 mg, 1.55 mmol).

¹H-NMR (CDCl₃) δ: 1.41 (1H, m), 1.94 (2H, m), 2.26 (6H, s), 2.26-2.42 (3H, m), 2.53 (3H, s), 2.94 (3H, m), 7.09 (1H, d, J=8.1 Hz), 7.29-7.36 (3H, m), 7.46 (1H, s), 7.56 (2H, d, J=8.4 Hz), 7.67 (2H, d, J=8.1 Hz), 7.78 (1H, m), 7.92 (2H, d, J=9.0 Hz).

Elemental analysis for C₂₇H₃₀N₂OS · 0.2H₂O

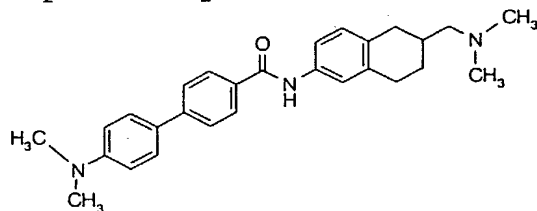
Calcd.: C, 74.69; H, 7.04; N, 6.45.

Found: C, 74.63; H, 7.03; N, 6.11.

Melting point: 178 - 180°C (crystallization solvent: ethyl acetate-diisopropyl ether)

Example 37

4'-(N,N-Dimethylamino)-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



4'-Amino-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide (150 mg, 0.375 mmol) obtained in Example 33, and paraformaldehyde (45.1 mg, 1.50 mmol) were suspended in mixed solution of methanol (1 ml) and tetrahydrofuran (1 ml). Sodium cyanohydroborate (94.4 mg, 1.50 mmol) was

added to the reaction mixture, which was stirred at 40°C for 18 hours. Ethyl acetate was added to the reaction mixture, which was washed with saturated aqueous sodium chloride solution, dried using anhydrous magnesium sulfate, and the solvent was distilled out under reduced pressure. The residue was refined using alumina column chromatography (development solvent; ethyl acetate), and pulverized using isopropyl ether, to give the titled compound (13 mg) as a white powder.

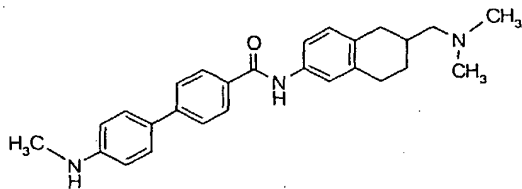
¹H-NMR (DMSO-d₆) δ: 1.32 (1H, m), 1.90 (2H, m), 2.15 (6H, s), 2.15-2.35 (3H, m), 2.77 (3H, m), 2.97 (6H, s), 6.82 (2H, d, J=8.4 Hz), 7.03 (1H, d, J=8.4 Hz), 7.48 (1H, d, J=8.1 Hz), 7.53 (1H, s), 7.63 (2H, d, J=8.7 Hz), 7.74 (2H, d, J=7.8 Hz), 7.98 (2H, d, J=8.4 Hz), 10.04 (1H, s).

FABMS(pos) 428.2[M+H]⁺

Melting point: 212 - 213°C (crystallization solvent: ethyl acetate-diisopropyl ether)

Example 38

N-[6-[(N,N-Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-4'-(methylamino)[1,1'-biphenyl]-4-carboxamide



The titled compound was obtained as a white powder by the same method as in Example 37, using 4'-amino-N-[6-[(N,N-dimethyl)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide (150 mg, 0.375 mmol) obtained in Example 33, paraformaldehyde (15.0 mg, 0.50 mmol), and sodium cyanohydroborate (31.5 mg, 0.50 mmol).

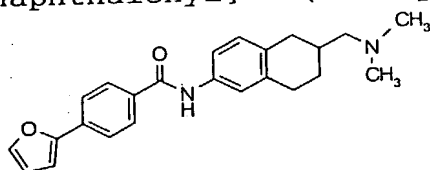
¹H-NMR (DMSO-d₆) δ: 1.32 (1H, m), 1.89 (2H, m), 2.15 (6H, s), 2.15-2.31 (3H, m), 2.72 (7H, m), 5.94 (1H, m), 6.64 (2H,

d, J=9.0 Hz), 7.03 (1H, d, J=8.7 Hz), 7.49 (4H, m), 7.70 (1H, d, J=8.4 Hz), 7.97 (2H, d, J=8.4 Hz), 10.02 (1H, s).
 FABMS(pos) 414.3[M+H]⁺
 Melting point: 163 - 165°C (crystallization solvent: ethyl acetate-diisopropyl ether)

5

Example 39

N-[6-[(N,N-Dimethyl)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-4-(2-furyl)benzamide



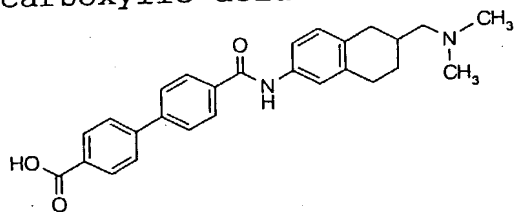
10

The titled compound (67 mg) was obtained as a white powder by the same method as in Example 16, using 4-bromo-N-[6-[(N,N-dimethyl)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]benzamide (250 mg, 0.645 mmol) obtained in Example 15, and 2-furylboric acid (86.7 mg, 0.775 mmol).
¹H-NMR (DMSO-d₆) δ: 1.40 (1H, m), 1.94 (2H, m), 2.25 (6H, s), 2.25-2.45 (3H, m), 2.88 (3H, m), 7.08 (1H, d, J=8.1 Hz), 7.26 (4H, m), 7.41 (1H, m), 7.60-7.74 (5H, m).
 FABMS(pos) 375.2[M+H]⁺

20

Example 40

4'-[[[6-[(N,N-Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]amino]carbonyl][1,1'-biphenyl]-4-carboxylic acid



25

Ethyl-4'-[[[6-[(N,N-dimethyl)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]amino]carbonyl][1,1'-biphenyl]-4-carboxylate (100 mg, 0.219 mmol) obtained in Example 35 was dissolved in a mixed solution of ethanol (3

ml) and water (0.5 ml). 1N aqueous sodium hydroxide solution (0.329 ml) was added to the reaction mixture at room temperature, which was stirred at 90°C for 5 hours.

After the solvent was distilled out under reduced
5 pressure, water was added to the residue, then 1N hydrochloric acid (0.329 ml) was added and the reaction mixture was stirred. The precipitated crude product collected by filtration, and washed with water to give the titled compound (89 mg) as a white powder.

10 ¹H-NMR (DMSO-d₆) δ: 1.34 (1H, m), 1.91 (2H, m), 2.24 (6H, s), 2.24-2.30 (3H, m), 2.81 (3H, m), 7.05 (1H, d, J=8.4 Hz), 7.49 (1H, d, J=8.4 Hz), 7.55 (1H, s), 7.89 (4H, m), 8.07 (4H, m), 10.18 (1H, s).

Elemental analysis for C₂₇H₂₈N₂O₃ · 2H₂O

15 Calcd.: C, 69.81; H, 6.94; N, 6.03.

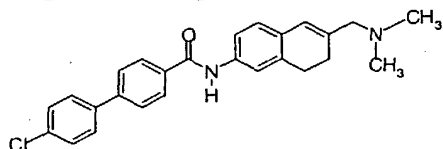
Found: C, 69.57; H, 7.01; N, 5.93.

Melting point: 143°C (decomposition) (crystallization solvent: water)

20

Example 41

4'-Chloro-N-[6-[(N,N-dimethyl)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



25

1) 6-Acetamido-1-tetralone (5.0 g, 0.0246 mol) synthesized according to a known method by documents (Journal of Organic Chemistry 27, 70 (1962)), was dissolved in 50 ml of DMF dimethylacetate, which was stirred at 110°C for 2 hours. The precipitate was collected by filtration,
30 and washed with ethyl acetate to give 6-acetamido-2-(N,N-dimethylaminomethylidene)-1-tetralone (4.98 g) as a yellow powder.

¹H-NMR (CDCl₃) δ: 2.19 (3H, s), 2.79-2.83 (2H, m), 2.88-

2.92 (2H, m), 3.11 (6H, s), 7.14-7.17 (1H, m), 7.68 (1H, s), 7.69 (1H, s), 7.95 (1H, d, J=8.1Hz), 7.96 (1H, s).

Melting point: 207 - 210°C (crystallization solvent: ethyl acetate)

5 2) The obtained 6-acetamido-2-(N,N-dimethylaminomethylidene)-1-tetralone (4.50 g, 0.0173 mol) was dissolved in methanol (50 ml), and sodium borohydride (6.56 g, 0.173 mol) was added to the solution under ice-cooling, which was stirred for 2 hours. The
10 reaction mixture was concentrated. Ethyl acetate and sodium hydrogencarbonate solution were added to the residue, and extraction was conducted. The ethyl acetate layer was concentrated, and 30 ml of tetrahydrofuran and 30 ml of 2N hydrochloric acid were added to the residue,
15 which was refluxed with heating for 16 hours. The reaction mixture was concentrated, and ethyl acetate and 2N sodium hydroxide solution were added, and extraction was conducted. The ethyl acetate layer was concentrated, and the residue was refined using alumina column chromatography
20 (development solvent; ethyl acetate:n-hexane = 30:70), to give 6-[(N,N-dimethylamino)methyl]-7,8-dihydro-2-naphthaleneamine (1.60 g) as a colorless oily substance.
¹H-NMR (CDCl₃) δ: 2.23 (6H, s), 2.28 (2H, t, J=8.4Hz), 2.74 (2H, t, J=8.4Hz), 2.95 (2H, s), 3.57-3.72 (2H, m), 6.25 (1H, s),
25 6.46-6.48 (2H, m), 6.83 (1H, d, J=8.7Hz).

3) The titled compound (1.12 g) was obtained as a white powder by the same method as in Example 1, using the obtained 6-[(N,N-dimethylamino)methyl]-7,8-dihydro-2-naphthalenamine (1.00 g, 0.005 mol), and 4-chlorobiphenyl
30 carboxylic acid (2.31 g, 0.01 mol).

¹H-NMR (CDCl₃) δ: 2.25 (6H, s), 2.34 (2H, t, J=7.8Hz), 2.86 (2H, t, J=7.8Hz), 2.99 (2H, s), 6.34 (1H, s), 7.03 (1H, d, J=8.7Hz), 7.39 (1H, d, J=8.1 Hz), 7.45 (2H, d, J=8.7), 7.48 (1H, s), 7.56 (2H, d, J=8.4 Hz), 7.67 (2H, d, J=8.4 Hz),
35 7.78 (1H, s), 7.94 (2H, d, J=8.4 Hz).

Elemental analysis for C₂₆H₂₅ClN₂O

Calcd.: C, 74.90; H, 6.04; N, 6.72.

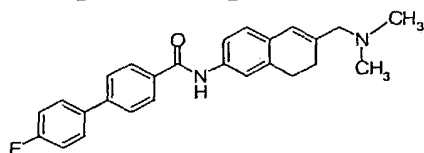
Found: C, 74.64; H, 6.14; N, 6.56.

Melting point: 204 - 207°C (crystallization solvent: ethyl acetate - n-hexane)

5

Example 42

4'-Fluoro-N-[6-[(N,N-dimethylamino)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



10 The titled compound (990 mg) was obtained as a white powder by the same method as in Example 1, using 6-[(N,N-dimethylamino)methyl]-7,8-dihydro-2-naphthalenamine (936 mg, 4.62 mmol) obtained in Example 41-2), and 4-fluorobiphenylcarboxylic acid (1.00 g, 4.62 mmol).

¹H-NMR (CDCl₃) δ: 2.25 (6H, s), 2.34 (2H, t, J=8.1Hz), 2.85 (2H, t, J=8.1Hz), 2.99 (2H, s), 6.34 (1H, s), 7.02 (1H, d, J=8.1Hz), 7.13-7.19 (2H, m), 7.38-7.41 (1H, m), 7.48 (1H, s), 7.56-7.61 (2H, m), 7.65 (2H, d, J=8.4 Hz), 7.80 (1H, s), 7.93 (2H, d, J=8.5Hz).

20

Elemental analysis for C₂₆H₂₅FN₂O

Calcd.: C, 77.97; H, 6.29; N, 6.99.

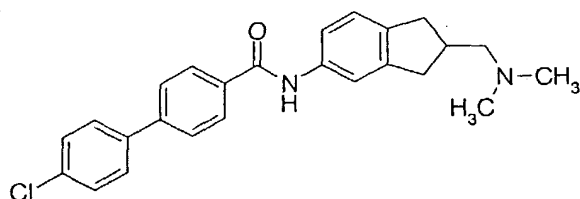
Found: C, 77.90; H, 6.23; N, 6.58.

Melting point: 190 - 193°C (crystallization solvent: ethyl acetate - n-hexane)

25

Example 43

4'-Chloro-N-[2-[(dimethylamino)methyl]-2,3-dihydro-1H-inden-5-yl][1,1'-biphenyl]-4-carboxamide



Concentrated hydrochloric acid (1 ml) was added to N-[2-[(dimethylamino)methyl]-2,3-dihydro-1H-inden-5-yl]acetamide (48.9 mg, 0.210 mmol) obtained in Reference
 5 Example 48, which was stirred at 110°C for 2 hours, and the solvent was distilled out under reduced pressure. Ethyl acetate was added to the residue, which was washed with potassium carbonate solution and saturated aqueous sodium chloride solution, dried using anhydrous sodium sulfate,
 10 and then the solvent was distilled out under reduced pressure. Using the oily substance obtained, the same operation as in Example 1 was conducted to give the titled compound (30 mg).

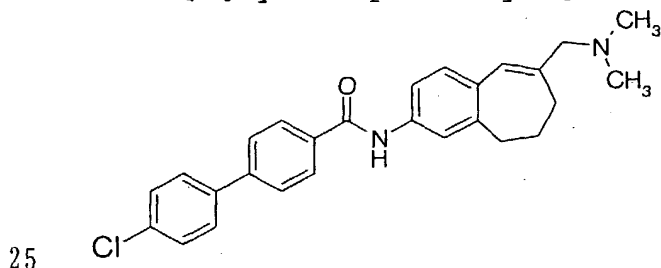
¹H NMR (DMSO-d₆) δ: 2.16 (6H, s), 2.22 (2H, d, J = 6.7 Hz),
 15 2.61 (4H, m), 2.97 (1H, m), 7.15 (1H, d, J = 8.1 Hz), 7.47 (1H, d, J = 8.1 Hz), 7.56 (2H, d, J = 8.4 Hz), 8.05 (2H, d, J = 8.4 Hz), 10.17 (1H, s).

FAB(pos) 405.1 [M+H]⁺

Melting point: 192 - 194°C (crystallization solvent: ethyl
 20 acetate - diisopropyl ether)

Example 44

4'-Chloro-N-[8-[(dimethylamino)methyl]-6,7-dihydro-5H-benzo[a]cyclohepten-3-yl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 8-

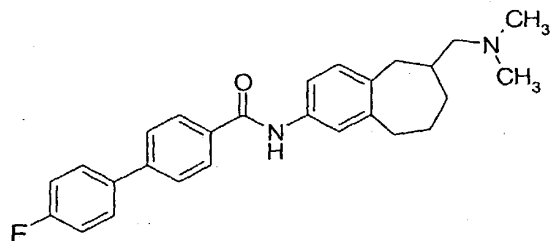
[(dimethylamino)methyl]-6,7-dihydro-5H-benzo[a]cyclohepten-3-amine obtained in Reference Example 50.

¹H-NMR (CDCl₃) δ: 1.96-2.10 (2H, m), 2.25 (6H, s), 2.39 (2H, t, J = 6.4 Hz), 2.79-2.85 (2H, m), 2.96 (2H, s), 6.40 (1H, s), 7.15 (1H, d, J = 8.6 Hz), 7.40-7.52 (4H, m), 7.56 (2H, d, J = 8.4 Hz), 7.67 (2H, d, J = 8.1 Hz), 7.81 (1H, s), 7.94 (2H, d, J = 8.1 Hz).

Melting point: 183-185°C (crystallization solvent: ethyl acetate - diethyl ether)

Example 45

4'-Fluoro-N-[6-[(dimethylamino)methyl]-6,7,8,9-tetrahydro-5H-benzo[a]cyclohepten-2-yl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 6-[(dimethylamino)methyl]-6,7,8,9-tetrahydro-5H-benzo[a]cyclohepten-2-amine obtained in Reference Example 51.

¹H-NMR (CDCl₃) δ: 1.40-1.68 (3H, m), 1.85-2.20 (10H, m), 2.55-2.92 (4H, m), 7.13-7.20 (3H, m), 7.35-7.43 (2H, m), 7.56-7.67 (4H, m), 7.77 (1H, s), 7.93 (2H, d, J=8.4 Hz).

Elemental analysis for C₂₇H₂₉FN₂O

Calcd.: C, 77.85; H, 7.02; N, 6.73.

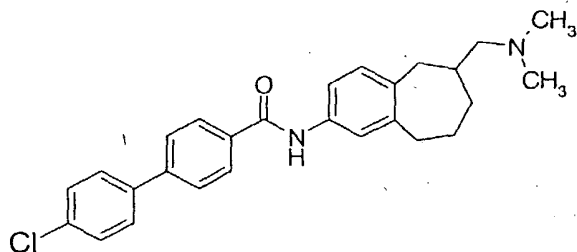
Found: C, 78.18; H, 7.09; N, 6.74.

Melting point: 167 - 169°C (crystallization solvent: diethyl ether)

Example 46

4'-Chloro-N-[6-[(dimethylamino)methyl]-6,7,8,9-

tetrahydro-5H-benzo[a]cyclohepten-2-yl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the same operation as in Experiment Example 1, using 6-[(dimethylamino)methyl]-6,7,8,9-tetrahydro-5H-benzo[a]cyclohepten-2-amine obtained in Reference Example 51.

$^1\text{H-NMR}$ (CDCl_3) δ : 1.40-1.67 (3H, m), 1.85-2.20 (10H, m), 2.55-2.92 (4H, m), 7.15 (1H, d, $J = 8.1$ Hz), 7.35-7.46 (4H, m), 7.56 (2H, d, $J = 8.4$ Hz), 7.66 (2H, d, $J = 8.1$ Hz), 7.77 (1H, s), 7.93 (2H, d, $J = 8.4$ Hz).

Elemental analysis for $\text{C}_{27}\text{H}_{29}\text{ClN}_2\text{O}$

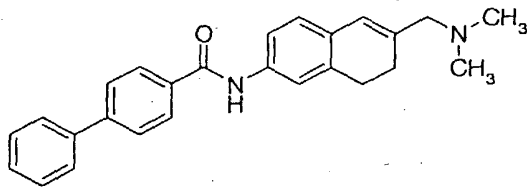
Calcd.: C, 74.90; H, 6.75; N, 6.47.

Found: C, 74.77; H, 6.65; N, 6.43.

Melting point: 173 - 175°C (crystallization solvent: diethyl ether)

Example 47

N-[6-[(Dimethylamino)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 6-[(N,N-dimethylamino)methyl]-7,8-dihydro-2-naphthalenamine obtained in Example 41-2).

$^1\text{H NMR}$ (CDCl_3) δ : 2.25 (6H, s), 2.33 (2H, t, $J = 5.4$ Hz),

2.84 (2H, t, $J = 5.4$ Hz), 2.98 (2H, s), 6.34 (1H, s), 7.01 (1H, d, $J = 7.8$ Hz), 7.32-7.94 (12H, m).

Elemental analysis for $C_{26}H_{26}N_2O$

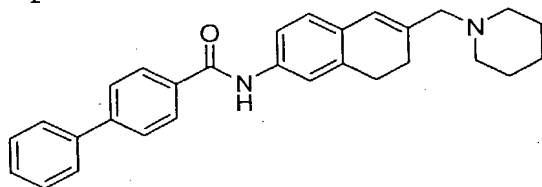
Calcd.: C, 81.64; H, 6.85; N, 7.32.

5 Found: C, 81.65; H, 6.79; N, 6.91.

Melting point: 173 - 175°C (crystallization solvent: tetrahydrofuran - n-hexane)

Example 48

10 N-[6-(1-Piperidinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



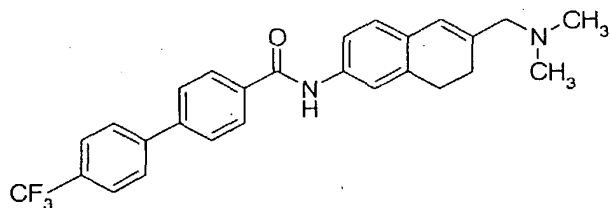
The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(1-piperidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 52.

1H NMR ($CDCl_3$) δ : 1.46-1.59 (6H, m), 2.31-2.36 (6H, m), 2.84 (2H, t, $J = 8.0$ Hz), 3.02 (2H, s), 6.34 (1H, s), 7.02 (1H, d, $J = 8.1$ Hz), 7.37-7.50 (4H, m), 7.63 (2H, d, $J = 6.9$ Hz), 7.71 (2H, d, $J = 8.1$ Hz), 7.79 (1H, s), 7.94 (2H, d, $J = 8.1$ Hz).

Melting point: 156 - 158°C (crystallization solvent: tetrahydrofuran - n-hexane)

25 Example 49

N-[6-[(Dimethylamino)methyl]-7,8-dihydro-2-naphthalenyl]-4'-trifluoromethyl[1,1'-biphenyl]-4-carboxamide



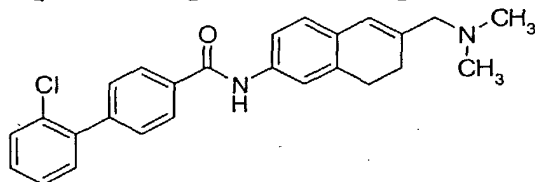
The titled compound was obtained by carrying out the same operation as in Example 1, using 6-[(N,N-dimethylamino)methyl]-7,8-dihydro-2-naphthalenamine obtained in Example 41-2).

^1H NMR (CDCl_3) δ : 2.25 (6H, s), 2.34 (d, $J = 5.1$ Hz), 2.86 (2H, d, $J = 5.1$ Hz), 2.99 (2H, s), 6.35 (1H, s), 7.04 (1H, d, $J = 8.4$ Hz), 7.40 (1H, d, $J = 3.3$ Hz), 7.49 (1H, s), 7.70-7.79 (6H, m), 7.87 (2H, d, $J = 8.4$ Hz).

Melting point: 214 - 216°C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 50

2'-Chloro-N-[6-[(dimethylamino)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 6-[(N,N-dimethylamino)methyl]-7,8-dihydro-2-naphthalenamine obtained in Example 41-2).

^1H NMR (CDCl_3) δ : 2.25 (6H, s), 2.34 (d, $J = 5.1$ Hz), 2.85 (2H, d, $J = 5.1$ Hz), 3.00 (2H, s), 6.34 (1H, s), 6.69 (1H, s), 7.02 (1H, d, $J = 8.4$ Hz), 7.31-7.57 (8H, m), 7.85 (1H, s), 7.92 (2H, d, $J = 7.8$ Hz).

Elemental analysis for $\text{C}_{26}\text{H}_{25}\text{ClN}_2\text{O}$

Calcd.: C, 74.90; H, 6.04; N, 6.72

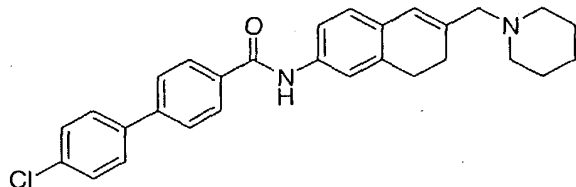
Found: C, 74.49; H, 5.65; N, 6.06.

Melting point: 145 - 147°C (crystallization solvent: ethyl

acetate - n-hexane)

Example 51

4'-Chloro-N-[6-(1-piperidinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



After N,N-dimethylformaldehyde solution (5 ml) of 4'-chloro-N-[6-(chloromethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide (225 mg) obtained in Reference Example 56, piperidine (0.16 ml), and diisopropylethylamine (0.282 ml) was stirred at room temperature for 15 hours, which was heated at 120° C for 2 hours. The residue obtained by concentrating the reaction mixture was dissolved in water-ethyl acetate, then extracted using ethyl acetate. The extract was washed with saturated aqueous sodium chloride solution, dried using anhydrous magnesium sulfate, and then the solvent was distilled out under reduced pressure. The resulting residue was refined using alumina column chromatography (development solvent; tetrahydrofuran:n-hexane = 1:5), and crystallized using tetrahydrofuran - n-hexane to give the titled compound (110 mg).

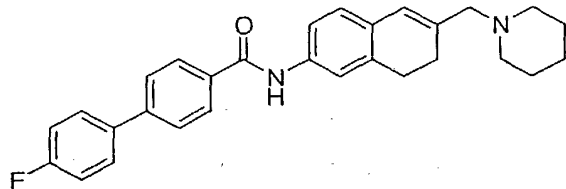
¹H NMR (CDCl₃) δ: 1.26-1.61 (6H, m), 2.30-2.36 (6H, m), 2.83 (2H, t, J = 8.4 Hz), 3.02 (2H, s), 6.33 (1H, s), 7.01 (1H, d, J = 8.1 Hz), 7.36-7.49 (4H, m), 7.55 (2H, d, J = 8.4 Hz), 7.66 (2H, d, J = 8.4 Hz), 7.81 (1H, s), 7.93 (2H, d, J = 8.1 Hz).

Melting point: 209 - 211° C (crystallization solvent: tetrahydrofuran - n-hexane)

Example 52

4'-Fluoro-N-[6-(1-piperidinylmethyl)-7,8-dihydro-2-

naphthalenyl][1,1'-biphenyl]4-carboxamide



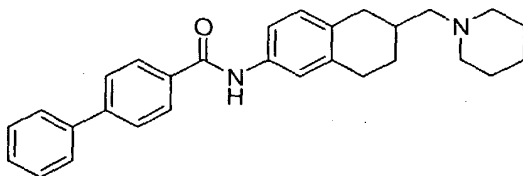
The titled compound was obtained by carrying out the same operation as in Example 1, using the 6-(1-piperidinylmethyl)-7,8-dihydro-2-naphthalene amine obtained in Reference example 52.

^1H NMR (CDCl_3) δ : 1.45-1.58 (6H, m), 2.29-2.37 (6H, m), 2.82 (2H, t, $J = 8.0$ Hz), 3.01 (2H, s), 6.33 (1H, s), 6.98-7.93 (12H, m).

Melting point: 190 - 192°C (crystallization solvent: tetrahydrofuran - n-hexane)

Example 53

N-[6-(1-Piperidinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(1-piperidinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenamine obtained in Reference Example 53.

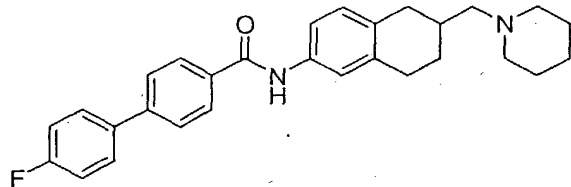
^1H NMR (CDCl_3) δ : 1.37-1.60 (8H, m), 1.96-2.00 (2H, m), 2.24-2.44 (5H, m), 2.82-2.93 (3H, m), 7.09 (1H, d, $J = 8.3$ Hz), 7.30-7.33 (1H, m), 7.38-7.65 (6H, m), 7.70 (2H, d, $J = 8.4$ Hz), 7.76 (1H, s), 7.93 (2H, d, $J = 8.4$ Hz).

Melting point: 160 - 162°C (crystallization solvent: tetrahydrofuran - n-hexane)

Example 54

4'-Fluoro-N-[6-[1-piperidinylmethyl)-5,6,7,8-

tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(1-piperidinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenamine obtained in Reference Example 53.

¹H NMR (CDCl₃) δ: 1.36-1.52 (8H, m), 2.29-2.31 (2H, m), 2.24-2.45 (6H, m), 2.82-2.93 (3H, m), 7.08-7.33 (4H, m), 7.44 (1H, s), 7.57-7.66 (4H, m), 7.74 (1H, s), 7.92 (2H, J = 8.1 Hz).

Elemental analysis for C₂₉H₃₁N₂O

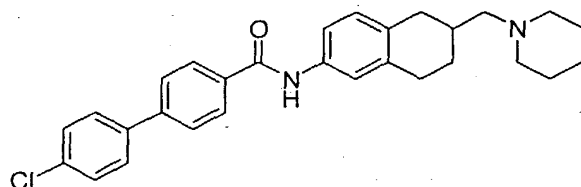
Calcd.: C, 78.70; H, 7.08; N, 6.33.

Found: C, 78.40; H, 7.09; N, 6.09.

Melting point: 179 - 181°C (crystallization solvent: ethyl acetate)

Example 55

4'-Chloro-N-[6-[1-piperidinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



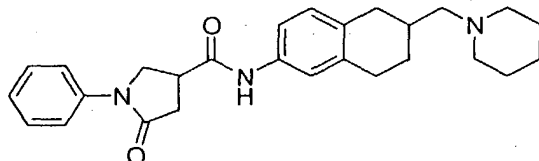
The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(1-piperidinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenamine obtained in Reference Example 53.

¹H NMR (CDCl₃) δ: 1.25-1.71 (8H, m), 1.95-2.00 (2H, m), 2.25-2.45 (6H, m), 2.83-2.93 (3H, m), 7.09 (1H, d, J = 8.3 Hz), 7.30-7.32 (1H, m), 7.43-7.45 (3H, m), 7.55 (2H, d, J = 8.1 Hz), 7.65 (2H, d, J = 8.4 Hz), 7.77 (1H, s), 7.93 (2H, d, J = 8.1 Hz).

Melting point: 202 - 203°C (crystallization solvent:
tetrahydrofuran - n-hexane)

Example 56

- 5 5-Oxo-1-phenyl-N-[6-(1-piperidinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenyl]-3-pyrrolidinecarboxamide



- The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(1-piperidinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenamine obtained in Reference Example 53.

^1H NMR (CDCl_3) δ : 1.03-3.33 (22H, m), 3.97 (1H, t, $J = 8.4$ Hz), 4.21 (1H, dd, $J = 6.8, 7.1$ Hz), 6.91-7.63 (9H, m).

Elemental analysis for $\text{C}_{27}\text{H}_{33}\text{N}_3\text{O}_2$

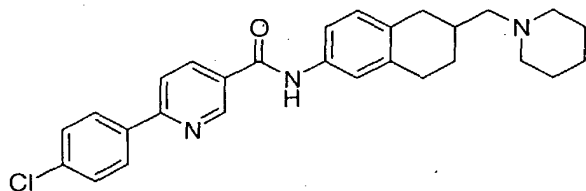
- 15 Calcd.: C, 75.14; H, 7.71; N, 9.74.

Found: C, 75.01; H, 7.33; N, 9.43.

Melting point: 162 - 164°C (crystallization solvent: ethyl acetate)

- 20 Example 57

6-(4-Chlorophenyl)-N-[6-(1-piperidinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenyl]nicotinamide



- The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(1-piperidinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenamine obtained in Reference Example 53.

^1H NMR (CDCl_3) δ : 1.30-2.40 (16H, m), 2.82-2.92 (3H, m), 7.09 (1H, d, $J = 8.1$ Hz), 7.26-7.48 (4H, m), 7.80 (2H, d,

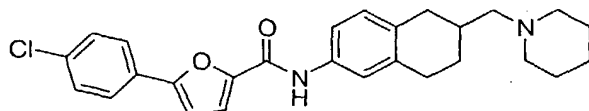
$J = 8.7$ Hz), 7.99 (2H, d, $J = 8.7$ Hz), 8.23 (d, 1H, $J = 6.3$ Hz), 9.11 (1H, s).

Melting point: 193 - 195°C (crystallization solvent: ethyl acetate)

5

Example 58

5-(4-Chlorophenyl)-N-[6-(1-piperidinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenyl]-2-furamide



10

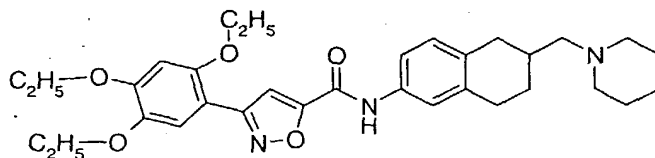
The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(1-piperidinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenamine obtained in Reference Example 53.

^1H NMR (CDCl_3) δ : 1.23-1.61 (7H, m), 1.96-2.00 (2H, m), 2.24-2.43 (7H, m), 2.80-2.92 (3H, m), 6.75 (1H, d, $J = 3.6$ Hz), 7.07 (1H, d, $J = 8.4$ Hz), 7.27 (1H, d, $J = 3.6$ Hz), 7.32-7.42 (4H, m), 7.66 (2H, d, $J = 8.4$ Hz), 8.32 (1H, s).

15

Example 59

20 N-[6-(1-Piperidinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenyl]-3-(2,4,5-triethoxyphenyl)-5-isoxazolecarboxamide



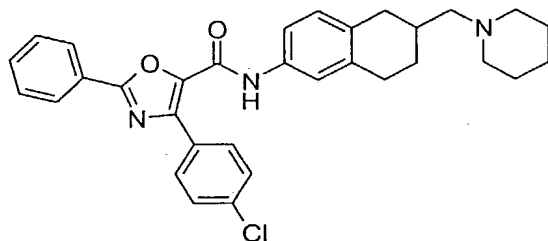
25 The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(1-piperidinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenamine obtained in Reference Example 53.

^1H NMR (CDCl_3) δ : 1.42-1.60 (18H, m), 1.97-2.36 (7H, m), 2.80-2.95 (3H, m), 4.06-4.18 (6H, m), 6.58 (1H, s), 7.09 (1H, d, $J = 8.4$ Hz), 7.35 (1H, d, $J = 8.1$ Hz), 7.44 (1H, s), 7.50 (1H, s), 7.55 (1H, s), 8.16 (1H, s).

30

Example 60

4-(4-Chlorophenyl)-2-phenyl-N-[6-(1-piperidinylmethyl)-
5,6,7,8-tetrahydro-2-naphthalenyl]-1,3-oxazole-5-
5 carboxamide

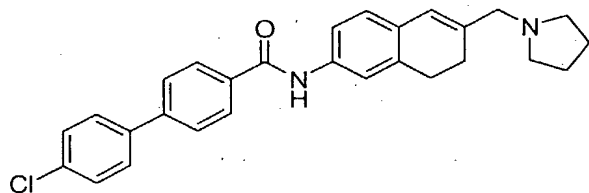


The titled compound was obtained by carrying out the
same operation as in Example 1, using 6-(1-
piperidinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenamine
10 obtained in Reference Example 53.

^1H NMR (CDCl_3) δ : 1.26-1.58 (7H, m), 1.90-2.00 (2H, m),
2.22-2.35 (7H, m), 2.70-2.95 (3H, m), 7.06 (1H, d, J = 8.1
Hz), 7.25-7.51 (7H, m), 8.04-8.32 (5H, m).

15 Example 61

4'-Chloro-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-
naphthalenyl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the
20 same operation as in Example 51, using 4'-chloro-N-[6-
(chloromethyl)-7,8-dihydro-2-naphthalenyl][1,1'-
biphenyl]-4-carboxamide obtained in Reference Example 56.
Melting point: 185 - 187°C (crystallization solvent:

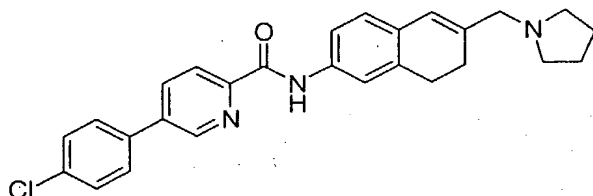
tetrahydrofuran - n-hexane)

25 ^1H NMR (CDCl_3) δ : 1.83 (4H, s), 2.35 (2H, t, J = 8.1 Hz),
2.52 (4H, s), 2.84 (2H, t, J = 8.1 Hz), 3.18 (2H, s), 6.36
(1H, s), 7.02 (1H, d, J = 8.4 Hz), 7.39-7.56 (6H, m), 7.66

(2H, d, $J = 7.5$ Hz), 7.82 (1H, s), 7.93 (2H, d, $J = 7.5$ Hz).

Example 62

5-(4-Chlorophenyl)-N-[6-(1-pyrrolidinymethyl)-7,8-dihydro-2-naphthalenyl]-2-pyridinecarboxamide

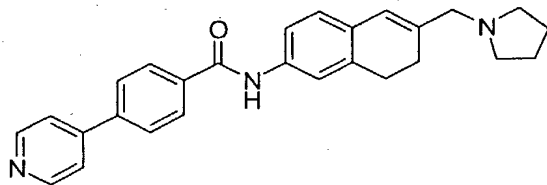


The titled compound was obtained by carrying out the same operation as in Example 51, using 4'-chloro-N-[6-(chloromethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide obtained in Reference Example 56.

^1H NMR (CDCl_3) δ : 1.80 (6H, s), 2.37 (2H, t, $J = 8.1$ Hz), 2.52 (4H, s), 2.87 (2H, t, $J = 8.1$ Hz), 3.18 (2H, s), 6.37 (1H, s), 7.03 (1H, d, $J = 7.8$ Hz), 7.48-7.61 (6H, m), 8.04 (1H, dd, $J = 8.1, 2.1$ Hz), 8.35 (1H, d, $J = 8.1$ Hz), 8.78 (1H, s), 9.95 (1H, s).

Example 63

4-(4-Pyridinyl)-N-[6-(1-pyrrolidinymethyl)-7,8-dihydro-2-naphthalenyl]benzamide

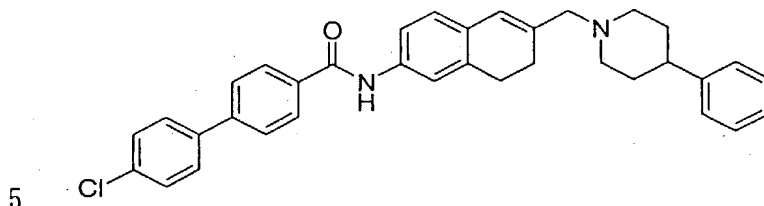


The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(1-pyrrolidinymethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

^1H NMR (CDCl_3) δ : 1.79-1.83 (6H, m), 2.35 (2H, t, $J = 8.1$ Hz), 2.53 (4H, s), 2.73 (2H, t, $J = 8.1$ Hz), 3.18 (2H, s), 6.36 (1H, s), 7.02 (1H, d, $J = 7.8$ Hz), 7.38 (1H, d, $J = 8.1$ Hz), 7.48 (1H, s), 7.71-7.78 (4H, m), 7.89 (1H, s), 7.99 (1H, d, $J = 8.4$ Hz), 8.32 (2H, d, $J = 8.4$ Hz).

Example 64

4'-Chloro-N-[6-[(4-phenyl-1-piperidinyl)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



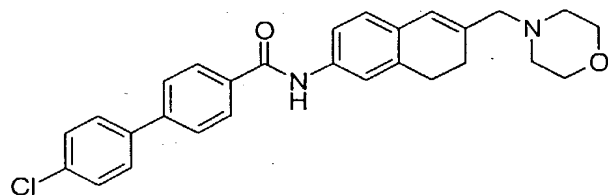
The titled compound was obtained by carrying out the same operation as in Example 51, using 4'-chloro-N-[6-(chloromethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide obtained in Reference Example 56.

10 ^1H NMR (CDCl_3) δ : 1.83-2.10 (6H, m), 2.37 (2H, t, $J = 8.1$ Hz), 2.47-2.54 (1H, m), 2.86 (2H, t, $J = 8.1$ Hz), 3.03-3.10 (2H, m), 3.10 (2H, s), 6.37 (1H, s), 7.03 (1H, d, $J = 8.4$ Hz), 7.19-7.57 (11H, m), 7.66 (2H, d, $J = 8.4$ Hz), 7.81 (1H, s), 7.94 (2H, d, $J = 8.4$ Hz).

15 Melting point: 228 - 230°C (crystallization solvent: tetrahydrofuran - n-hexane)

Example 65

20 4'-Chloro-N-[6-(4-morpholinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the same operation as in Example 51, using 4'-chloro-N-[6-(chloromethyl)-7,8-dihydro-2-naphthalenyl][1,1'-

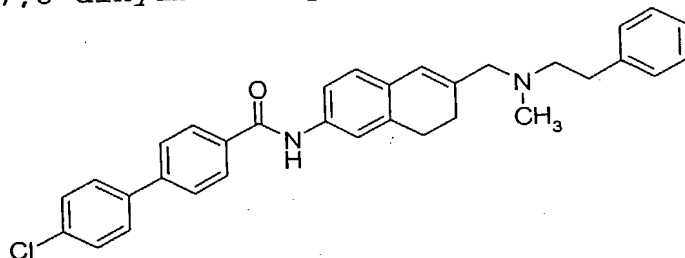
25 biphenyl]-4-carboxamide obtained in Reference Example 56.

^1H NMR (CDCl_3) δ : 2.34 (2H, t, $J = 7.8$ Hz), 2.45 (4H, s), 2.84 (2H, t, $J = 7.8$ Hz), 3.06 (2H, s), 3.73 (4H, s), 6.36 (1H, s), 7.02 (1H, d, $J = 8.1$ Hz), 7.36-7.57 (6H, m), 7.67 (2H, d, $J = 8.4$ Hz), 7.80 (1H, s), 7.94 (2H, d, $J = 8.4$ Hz).

Melting point: 194 - 195°C (crystallization solvent:
tetrahydrofuran - n-hexane)

Example 66

- 5 4'-Chloro-N-(6-[[methyl(2-phenylethyl)amino]methyl]-
7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide

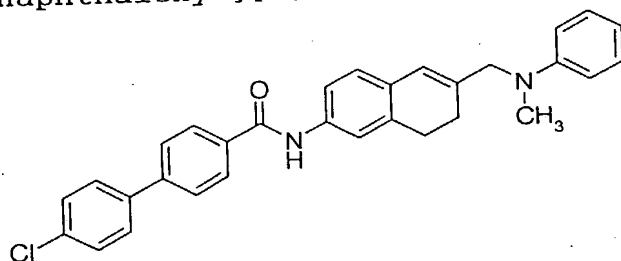


- The titled compound was obtained by carrying out the
same operation as in Example 51, using 4'-chloro-N-[6-
10 (chloromethyl)-7,8-dihydro-2-naphthalenyl][1,1'-
biphenyl]-4-carboxamide obtained in Reference Example 56.
¹H NMR (CDCl₃) δ: 2.25-2.32 (2H, m), 2.32 (3H, s), 2.60-2.66
(2H, m), 2.77-2.83 (4H, m), 3.10 (2H, s), 6.32 (1H, s),
6.93-7.95 (16H, m).

- 15 Melting point: 173 - 175°C (crystallization solvent:
tetrahydrofuran - n-hexane)

Example 67

- 20 4'-Chloro-N-[6-[methylanilino]methyl]-7,8-dihydro-2-
naphthalenyl][1,1'-biphenyl]-4-carboxamide



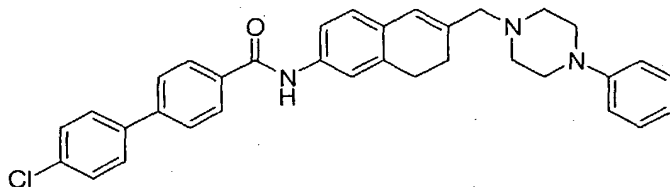
- The titled compound was obtained by carrying out the
same operation as in Example 51, using 4'-chloro-N-[6-
(chloromethyl)-7,8-dihydro-2-naphthalenyl][1,1'-
25 biphenyl]-4-carboxamide obtained in Reference Example 56.
¹H NMR (CDCl₃) δ: 2.20-2.30 (2H, m), 2.25 (3H, s), 2.85-2.90

(2H, m), 3.00 (2H, s), 6.30 (1H, s), 6.74-7.95 (146H, m).

Melting point: 177 - 179°C (crystallization solvent:
tetrahydrofuran - n-hexane)

5 Example 68

4'-Chloro-N-[6-[(4-phenyl-1-piperadiny)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide

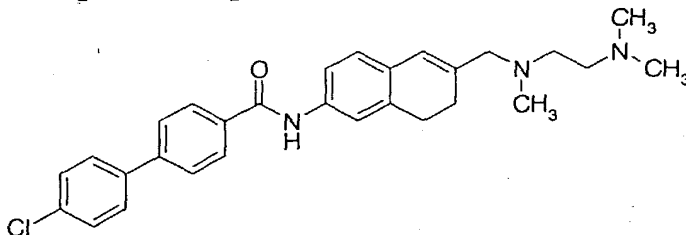


10 The titled compound was obtained by carrying out the same operation as in Example 51, using 4'-chloro-N-[6-(chloromethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide obtained in Reference Example 56.
¹H NMR (CDCl₃) δ: 2.37 (2H, t, J = 8.1 Hz), 2.62 (4h, S), 2.86 (2H, t, J = 8.4 Hz), 3.13 (2H, s), 3.22 (4H, s), 6.39
 15 (1H, s), 6.85-7.95 (16H, m).

Melting point: 228 - 230°C (crystallization solvent:
tetrahydrofuran - n-hexane)

Example 69

20 4'-Chloro-N-[6-[[[2-(dimethylamino)ethyl](methyl)amino]methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



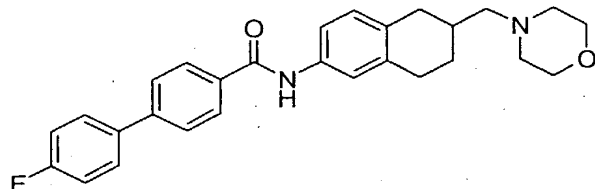
25 The titled compound was obtained by carrying out the same operation as in Example 51, using 4'-chloro-N-[6-(chloromethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide obtained in Reference Example 56.
¹H NMR (CDCl₃) δ: 2.25 (6H, s), 2.26 (3H, s), 2.33 (2H, t,

J = 8.1 Hz), 2.44-2.50 (4H, m), 2.84 (2H, t, J = 8.1 Hz), 3.07 (2H, s), 6.35 (1H, s), 7.02 (1H, d, J = 8.4 Hz), 7.37-7.57 (6H, m), 7.67 (2H, d, J = 8.1 Hz), 7.80 (1H, s), 7.94 (2H, d, J = 8.4 Hz).

- 5 Melting point: 156 - 158°C (crystallization solvent: tetrahydrofuran - n-hexane)

Example 70

- 10 4'-Fluoro-N-[6-(4-morpholinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



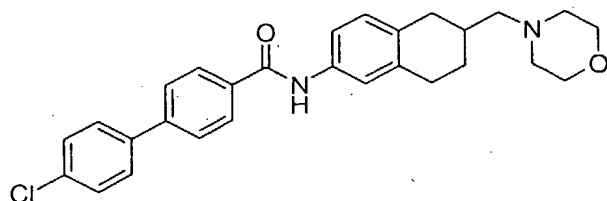
- 15 The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(4-morpholinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenamine obtained in Reference Example 57.

¹H NMR (CDCl₃) δ: 1.40-1.50 (1H, m), 1.90-2.10 (2H, m), 2.29-2.45 (7H, m), 2.80-2.92 (3H, m), 3.72-3.75 (4H, m), 7.07-7.33 (4H, m), 7.46 (1H, s), 7.56-7.66 (4H, m), 7.78 (1H, s), 7.92 (2H, d, J = 8.1 Hz).

- 20 Melting point: 188 - 190°C (crystallization solvent: ethyl acetate)

Example 71

- 25 4'-Chloro-N-[6-(4-morpholinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(4-morpholinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenamine

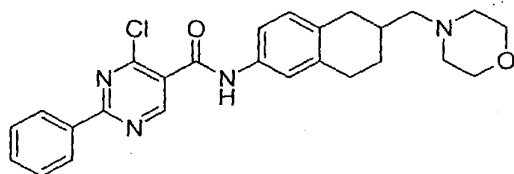
obtained in Reference Example 57.

^1H NMR (CDCl_3) δ : 1.40-1.50 (1H, m), 1.90-2.10 (2H, m), 2.32-2.45 (7H, m), 2.80-2.90 (3H, m), 3.70-3.80 (4H, m), 7.10-7.92 (12H, m).

5 Melting point: 216 - 218°C (crystallization solvent: ethyl acetate)

Example 72

10 4-Chloro-N-[6-(4-morpholinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenyl]-2-phenyl-5-pyrimidinecarboxamide

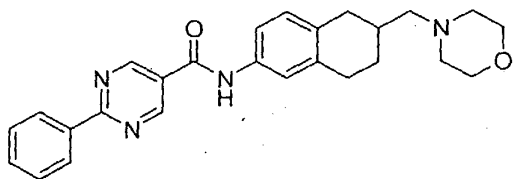


The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(4-morpholinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenamine
15 obtained in Reference Example 57.

^1H NMR (CDCl_3) δ : 1.40-1.50 (1H, m), 1.95-2.05 (2H, m), 2.29-2.45 (7H, m), 2.80-2.95 (3H, m), 3.73 (4H, t, J = 4.5 Hz), 7.10 (1H, d, J = 8.1 Hz), 7.32 (1H, d, J = 8.1 Hz), 7.42 (1H, s), 7.49-7.56 (3H, m), 8.25 (1H, s), 8.48 (2H, d, J = 6.6 Hz), 9.20 (1H, s)
20

Example 73

N-[6-(4-Morpholinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenyl]-2-phenyl-5-pyrimidinecarboxamide



25

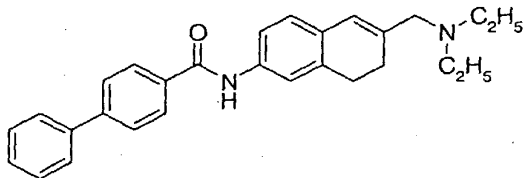
The titled compound was obtained by carrying out the same operation as in Reference Example 48, using 4-chloro-N-[6-(4-morpholinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenyl]-2-phenyl-5-pyrimidinecarboxamide obtained in

Example 72.

¹H NMR (CDCl₃) δ: 1.21-1.30 (1H, m), 1.93-2.03 (2H, m), 2.28-2.44 (7H, m), 2.80-2.90 (3H, m), 3.73 (4H, t, J = 4.8 Hz), 7.07 (1H, d, J = 8.1 Hz), 7.26-7.30 (1H, m), 7.39 (1H, s), 7.51-7.53 (3H, m), 8.00 (1H, s), 8.50 (2H, dd, J = 8.1, 2.4 Hz), 9.21 (2H, s)

Example 74

N-[6-[(Diethylamino)methyl]-7,8-dihydro-2-naphthalenyl]
[1,1'-biphenyl]-4-carboxamide



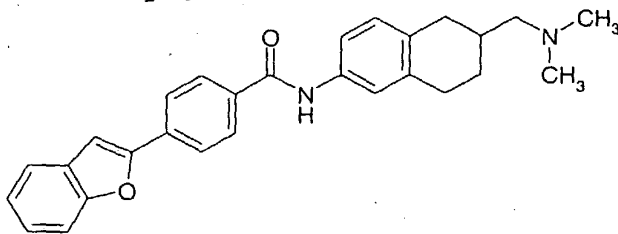
The titled compound was obtained by carrying out the same operation as in Example 51, using N-[6-(chloromethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide obtained in Reference Example 58.

¹H NMR (CDCl₃) δ: 1.24 (6H, t, J = 7.2 Hz), 2.33 (2H, t, J = 5.1 Hz), 2.53 (4H, q, J = 7.2 Hz), 2.84 (2H, t, J = 5.1 Hz), 3.11 (2H, s), 6.36 (1H, s), 7.02 (1H, d, J = 8.1 Hz), 7.37-7.50 (5H, m), 7.63 (2H, d, J = 8.7 Hz), 7.71 (2H, d, J = 8.4 Hz), 7.79 (1H, s), 7.93 (2H, d, J = 8.4 Hz).

Melting point: 153 - 155°C (crystallization solvent: tetrahydrofuran - n-hexane)

Example 75

4-(2-Benzo[b]furanyl)-N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]benzamide



The titled compound was obtained by carrying out the

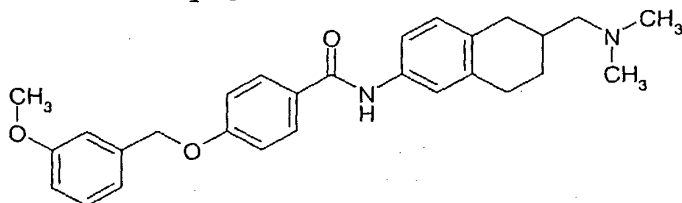
same operation as in Example 4, using 6-amino-2-[(N,N-dimethylamino)methyl]tetralin hydrochloride.

Melting point: 192 - 194°C (crystallization solvent: tetrahydrofuran-isopropyl ether)

5

Example 76

4-(3-Methoxybenzyloxy)-N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]benzamide



10

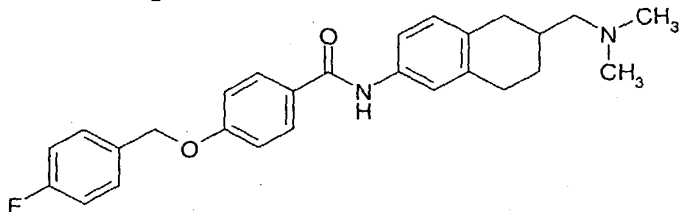
The titled compound was obtained by carrying out the same operation as in Example 4, using 6-amino-2-[(N,N-dimethylamino)methyl]tetralin hydrochloride.

Melting point: 102 - 104°C (crystallization solvent: isopropyl ether)

15

Example 77

4-(4-Fluorobenzyloxy)-N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]benzamide



20

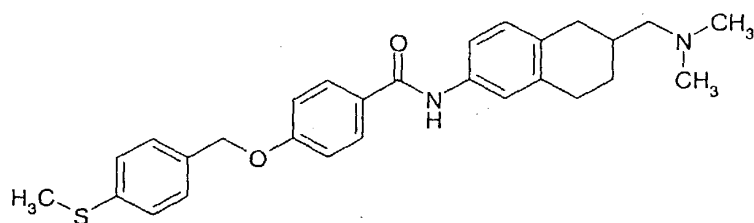
The titled compound was obtained by carrying out the same operation as in Example 4, using 6-amino-2-[(N,N-dimethylamino)methyl]tetralin hydrochloride.

Melting point: 165 - 167°C (crystallization solvent: tetrahydrofuran-hexane)

25

Example 78

4-[4-(Methylsulfanyl)benzyloxy]-N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]benzamide

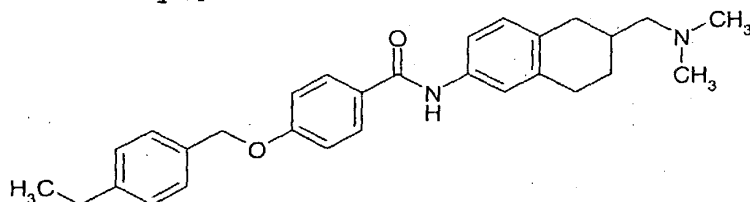


The titled compound was obtained by carrying out the same operation as in Example 4, using 6-amino-2-[(N,N-dimethylamino)methyl]tetralin hydrochloride.

- 5 Melting point: 162 - 163°C (crystallization solvent: tetrahydrofuran-hexane)

Example 79

- 4-(4-Ethylbenzyloxy)-N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]benzamide

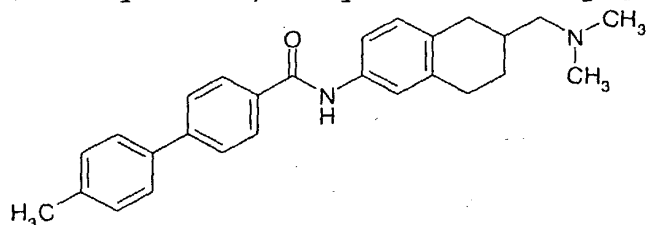


The titled compound was obtained by carrying out the same operation as in Example 4, using 6-amino-2-[(N,N-dimethylamino)methyl]tetralin hydrochloride.

- 15 Melting point: 120 - 122°C (crystallization solvent: tetrahydrofuran-isopropyl ether)

Example 80

- 20 (4'-Methylbiphenyl-4-yl)-N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]carboxamide



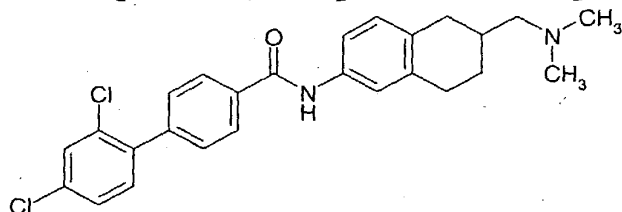
The titled compound was obtained by carrying out the same operation as in Example 4, using 6-amino-2-[(N,N-

dimethylamino)methyl]tetralin hydrochloride.

Melting point: 181 - 182°C (crystallization solvent: ethyl acetate-hexane)

5 Example 81

(2',4'-Dichlorobiphenyl-4-yl)-N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]carboxamide

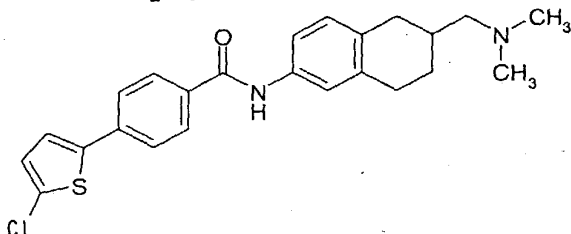


10 The titled compound was obtained by carrying out the same operation as in Example 4, using 6-amino-2-[(N,N-dimethylamino)methyl]tetralin hydrochloride.

Melting point: 188 - 189°C (crystallization solvent: tetrahydrofuran-hexane)

15 Example 82

4-(5-Chloro-2-thienyl-N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]benzamide

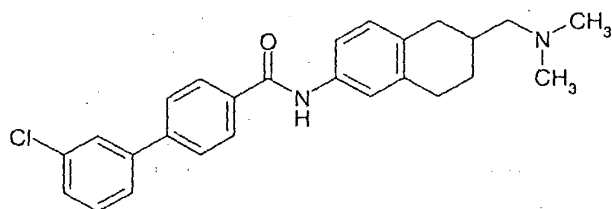


20 The titled compound was obtained by carrying out the same operation as in Example 4, using 6-amino-2-(N,N-dimethylamino)methyltetraline hydrochloride.

Melting point: 167 - 169°C (crystallization solvent: ethyl acetate-hexane)

25 Example 83

(3'-Chlorobiphenyl-4-yl)-N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]carboxamide

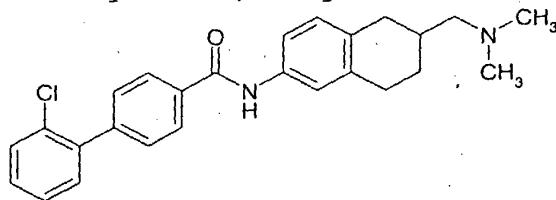


The titled compound was obtained by carrying out the same operation as in Example 4, using 6-amino-2-[(N,N-dimethylamino)methyl]tetralin hydrochloride.

5 Melting point: 138 - 139°C (crystallization solvent: tetrahydrofuran-isopropyl ether)

Example 84

(2'-Chlorobiphenyl-4-yl)-N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]carboxamide

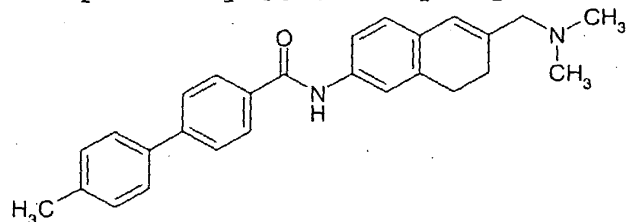


The titled compound was obtained by carrying out the same operation as in Example 4, using 6-amino-2-[(N,N-dimethylamino)methyl]tetralin hydrochloride.

15 Melting point: 176 - 177°C (crystallization solvent: tetrahydrofuran-hexane)

Example 85

4'-Methyl-N-[6-[N,N-dimethylamino)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 6-[(N,N-dimethylamino)methyl]-7,8-dihydro-2-naphthalenamine

obtained in Example 41-2).

¹H-NMR (CDCl₃) δ: 2.25 (6H, s), 2.33 (2H, t, J = 8.1 Hz),
2.41 (3H, s), 2.84 (2H, t, J = 8.1 Hz), 2.98 (2H, s), 6.33
(1H, s), 7.01 (1H, d, J = 7.8 Hz), 7.39 (1H, d, J = 8.4 Hz),
5 7.48 (1H, s), 7.52 (2H, d, J = 7.8 Hz), 7.67 (2H, d, J =
8.1 Hz), 7.84 (1H, s), 7.91 (2H, d, J = 8.1 Hz).

Elemental analysis for C₂₇H₂₈N₂O

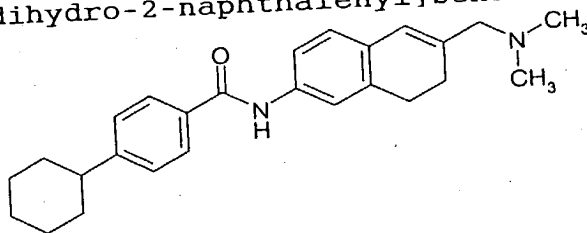
Calcd.: C, 81.78; H, 7.12; N, 7.06

Found: C, 81.51; H, 7.22; N, 6.93

10 Melting point: 195 - 196°C (crystallization solvent: ethyl
acetate-diisopropyl ether)

Example 86

15 4-Cyclohexyl-N-[6-[(N,N-dimethylamino)methyl]-7,8-
dihydro-2-naphthalenyl]benzamide



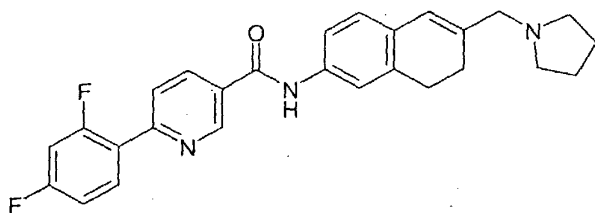
The titled compound was obtained by carrying out the
same operation as in Example 1, using the 6-[(N,N-
dimethylamino)methyl]-7,8-dihydro-2-naphthalenamine
20 obtained in Example 41-2).

¹H-NMR (CDCl₃) δ: 1.20-1.52 (4H, m), 1.71-1.96 (6H, m), 2.25
(6H, s), 2.33 (2H, t, J = 8.1 Hz), 2.50-2.62 (1H, m), 2.84
(2H, t, J = 8.1 Hz), 2.99 (2H, s), 6.33 (1H, s), 7.00 (1H,
d, J = 7.8 Hz), 7.31 (2H, d, J = 8.1 Hz), 7.36 (1H, d, J
25 = 7.8 Hz), 7.46 (1H, brs), 7.75 (1H, s), 7.78 (2H, d, J
= 8.1 Hz).

Melting point: 179 - 181°C (crystallization solvent: ethyl
acetate-diisopropyl ether)

30 Example 87

6-(2,4-Difluorophenyl)-N-[6-[(1-pyrrolidinyl)methyl]-
7,8-dihydro-2-naphthalenyl]nicotinamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

¹H-NMR (CDCl₃) δ: 1.81 (4H, m), 2.37 (2H, t, J = 8.1 Hz), 2.54 (4H, m), 2.86 (2H, t, J = 8.1 Hz), 3.18 (2H, s), 6.37 (1H, s), 6.93 (1H, m), 7.04 (2H, m), 7.38 (1H, m), 7.47 (1H, s), 7.77 (1H, s), 7.91 (1H, m), 8.13 (1H, m), 8.24 (1H, m), 9.16 (1H, s).

Elemental analysis for C₂₇H₂₆F₂N₃O

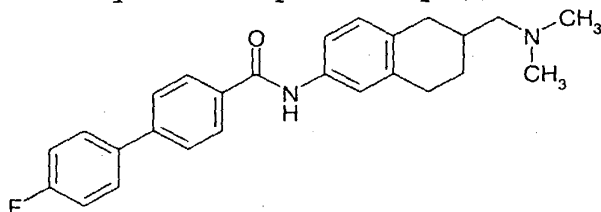
Calcd.: C, 72.79; H, 5.66; N, 9.43

Found: C, 72.65; H, 5.52; N, 9.73

Melting point: 169 - 170°C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 88

4'-Fluoro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the same operation as in Example 4, using 6-amino-2-[(N,N-dimethylamino)methyl]tetralin hydrochloride.

¹H-NMR (CDCl₃) δ: 1.41 (1H, m), 1.95 (2H, m), 2.25-2.45 (3H, m), 2.36 (6H, s), 2.85-2.94 (3H, m), 7.13 (3H, m), 7.30 (1H, m), 7.46 (1H, s), 7.59 (2H, m), 7.65 (2H, d, J = 8.1 Hz), 7.74 (1H, s), 7.93 (2H, d, J = 8.1 Hz).

Elemental analysis for C₂₆H₂₇FN₂O

Calcd.: C, 77.58; H, 6.76; N, 6.96

Found: C, 77.72; H, 6.49; N, 6.79

Melting point: 184 - 186°C (crystallization solvent: ethyl acetate - diisopropyl ether)

5

Example 89

(+)-4'-Fluoro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide, and (-)-4'-fluoro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide

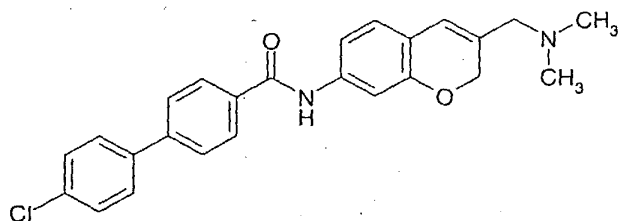
Optical resolution of 4'-fluoro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide (2.00 g) obtained in Example 88 was conducted by sample-splitting HPLC using a chiral column (Daicel Co., CHIRALCEL OD 500 mmD x 500 mmL; moving phase n-hexane:ethanol = 85:15), to give (+) form (1.00 g; 99.8%ee) and (-) form (0.89 g; >99.9%ee) as powders. The powders obtained were respectively recrystallized using ethyl acetate - diisopropyl ether, to give the (+) form (855 mg) and (-) form (754 mg) of the titled compounds. The optical rotation of both compounds are shown below.

(+)-4'-fluoro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide
Optical rotation: $[\alpha]_D = +50.8^\circ$ C=0.494% (methanol)
(-)-4'-fluoro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide
Optical rotation: $[\alpha]_D = +51.2^\circ$ C=0.492% (methanol)

30

Example 90

4'-Chloro-N-[3-[(N,N-dimethylamino)methyl]-2H-chromen-7-yl][1,1'-biphenyl]-4-carboxamide



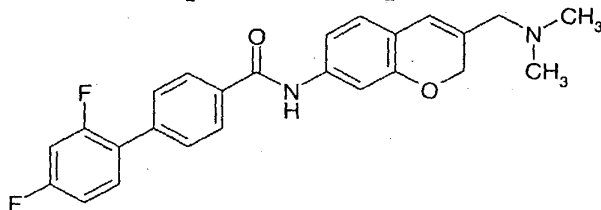
The titled compound was obtained by carrying out the same operation as in Example 1, using 3-[(N,N-dimethylamino)methyl]-2H-chromen-7-amine obtained in Reference Example 59.

¹H-NMR (CDCl₃) δ: 2.23 (6H, s), 2.97 (2H, s), 4.79 (2H, s), 6.30 (1H, s), 6.96 (1H, d, J = 8.1 Hz), 7.13 (1H, s), 7.20 (1H, d, J = 8.1 Hz), 7.45 (2H, d, J = 8.6 Hz), 7.56 (2H, d, J = 8.6 Hz), 7.66 (2H, d, J = 8.4 Hz), 7.74 (1H, brs), 7.93 (2H, d, J = 8.4 Hz).

Melting point: 199 - 208°C (crystallization solvent: diisopropyl ether)

Example 91

2',4'-Difluoro-N-[3-[(N,N-dimethylamino)methyl]-2H-chromen-7-yl][1,1'-biphenyl]-4-carboxamide



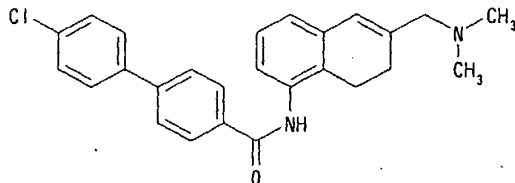
The titled compound was obtained by carrying out the same operation as in Example 1, using 3-[(N,N-dimethylamino)methyl]-2H-chromen-7-amine obtained in Reference Example 59.

¹H-NMR (CDCl₃) δ: 2.23 (6H, s), 2.97 (2H, s), 4.78 (2H, s), 6.29 (1H, s), 6.80-7.10 (2H, m), 6.96 (1H, d, J = 8.1 Hz), 7.13 (1H, s), 7.20 (1H, d, J = 8.1 Hz), 7.40-7.50 (1H, m), 7.62 (2H, d, J = 8.4 Hz), 7.76 (1H, brs), 7.92 (2H, d, J = 8.4 Hz).

Melting point: 200 - 204°C (crystallization solvent: diisopropyl ether)

Example 92

4'-Chloro-N-[6-[(dimethylamino)methyl]-7,8-dihydro-1-naphthalenyl][1,1'-biphenyl]-4-carboxamide



5

The titled compound was obtained in the same manner as in Example 1, using 6-[(dimethylamino)methyl]-7,8-dihydro-1-naphthalenamine obtained in Reference Example

10

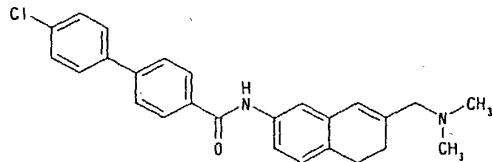
60. ¹H-NMR (CDCl₃) δ: 2.34 (6H, s), 2.36 (2H, t, J=8.1 Hz), 2.80 (2H, t, J=8.1 Hz), 3.00 (2H, s), 6.38 (1H, s), 6.94 (1H, d, J=7.8 Hz), 7.21 (1H, t, J=7.8 Hz), 7.45 (2H, d, J=8.6 Hz), 7.56 (2H, d, J=8.6 Hz), 7.61 (2H, m), 7.68 (2H, d, J=8.4 Hz), 7.97 (2H, d, J=8.4 Hz).

15

Melting point: 193 - 195°C (crystallization solvent : diisopropyl ether)

Example 93

20 4'-Chloro-N-[7-[(dimethylamino)methyl]-5,6-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained as a white powder by the same method as in Example 1, using 7-

25 [(dimethylamino)methyl]-5,6-dihydro-2-naphthalenamine obtained in Reference Example 61.

¹H-NMR (CDCl₃) δ: 2.25 (6H, s), 2.34 (2H, t, J=8.1 Hz), 2.82 (2H, t, J=8.1 Hz), 3.00 (2H, s), 6.36 (1H, s), 7.11 (1H, d, J=7.5 Hz), 7.34 (1H, d, J=8.1 Hz), 7.38 (1H, s), 7.44

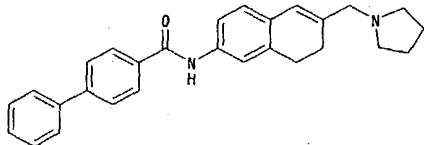
(2H, d, J=8.4 Hz), 7.56 (2H, d, J=8.4 Hz), 7.66 (2H, d, J=8.4 Hz), 7.78 (1H, brs), 7.97 (2H, d, J=8.4 Hz).

Melting point: 167 - 169°C (crystallization solvent : diisopropyl ether)

5

Example 94

N-[6-(1-Pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



10 The titled compound was obtained as a white powder in the same manner as in Example 1, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

¹H-NMR (CDCl₃) δ: 1.75-1.90 (4H, m), 2.34 (2H, t, J=8.1 Hz),
15 2.45-2.60 (4H, m), 2.85 (2H, t, J=8.1 Hz), 3.18 (2H, s),
6.36 (1H, s), 7.02 (1H, d, J=8.1 Hz), 7.27-7.55 (5H, m),
7.63 (2H, d, J=7.3 Hz), 7.70 (2H, d, J=8.4 Hz), 7.82 (1H, s),
7.94 (2H, d, J=8.1 Hz).

Elemental analysis for C₂₈H₂₈N₂O

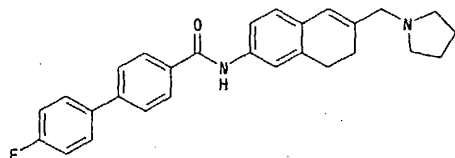
20 Calcd.: C, 82.32; H, 6.91; N, 6.86.

Found: C, 81.99; H, 6.69; N, 6.91.

Melting point: 176 - 177°C (crystallization solvent : diisopropyl ether)

25 Example 95

4'-Fluoro-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



30 The titled compound was obtained in the same manner as in Example 1, using 6-(1-pyrrolidinylmethyl)-7,8-

dihydro-2-naphthalenamine obtained in Reference Example 54.

¹H-NMR (CDCl₃) δ: 1.75-1.90 (4H, m), 2.35 (2H, t, J=8.2 Hz), 2.45-2.60 (4H, m), 2.84 (2H, t, J=8.2 Hz), 3.18 (2H, s), 5 6.36 (1H, s), 7.01 (1H, d, J=8.1 Hz), 7.16 (2H, t, J=8.1 Hz), 7.38 (1H, d, J=8.1 Hz), 7.48 (1H, brs), 7.56-7.61 (2H, m), 7.64 (2H, d, J=8.4 Hz), 7.83 (1H, s), 7.93 (2H, d, J=8.4 Hz).

Elemental analysis for C₂₈H₂₇N₂O

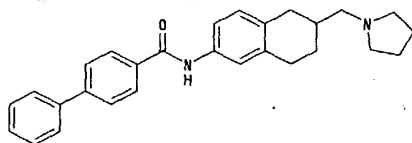
10 Calcd.: C, 78.85; H, 6.38; N, 6.57.

Found: C, 78.75; H, 6.39; N, 6.45.

Melting point: 189 - 192°C (crystallization solvent : diisopropyl ether)

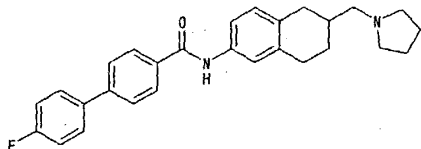
15 Example 96

N-[6-(1-Pyrrolidinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



Example 97

4'-Fluoro-N-[6-(1-pyrrolidinymethyl)-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



5 The titled compound was obtained as a white powder in the same manner as in Example 1, using 6-(1-pyrrolidinymethyl)-5,6,7,8-tetrahydro-2-naphthalenamine obtained in Reference Example 55.

10 ¹H-NMR (CDCl₃) δ: 1.40-1.50 (1H, m), 1.80 (4H, m), 1.80-2.10 (1H, m), 1.80-2.20 (8H, m), 3.30-4.00 (3H, m), 7.08 (1H, d, J=8.1 Hz), 7.15 (2H, t, J=8.4 Hz), 7.30 (1H, d, J=8.1 Hz), 7.44 (1H, brs), 7.56-7.61 (2H, m), 7.62 (2H, d, J=8.1 Hz), 7.85 (1H, s), 7.92 (2H, d, J=8.1 Hz).

Elemental analysis for C₂₈H₂₉FN₂O

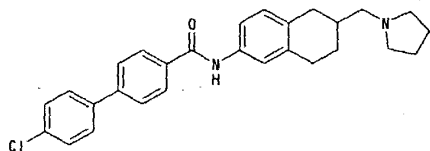
15 Calcd.: C, 78.48; H, 6.82; N, 6.54.

Found: C, 78.18; H, 6.60; N, 6.60.

Melting point: 185 - 189°C (crystallization solvent : diisopropyl ether)

20 Example 98

4'-Chloro-N-[6-(1-pyrrolidinymethyl)-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



25 The titled compound was obtained as a white powder in the same manner as in Example 1, using 6-(1-pyrrolidinymethyl)-5,6,7,8-tetrahydro-2-naphthalenamine obtained in Reference Example 55.

30 ¹H-NMR (CDCl₃) δ: 1.40-1.50 (1H, m), 1.80 (4H, m), 1.80-2.10 (1H, m), 1.80-2.20 (8H, m), 3.30-4.00 (3H, m), 7.08 (1H, d, J=8.1 Hz), 7.31 (1H, d, J=8.4 Hz), 7.43 (2H, d, J=8.7 Hz), 7.45 (1H, s), 7.54 (2H, d, J=8.7 Hz), 7.64 (2H, d, J=8.4

Hz), 7.80 (1H, s), 7.93 (2H, d, J=8.4 Hz).

Elemental analysis for $C_{28}H_{29}ClN_2O$

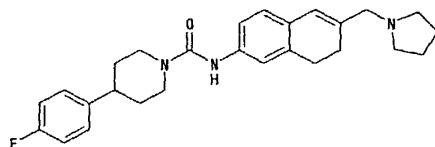
Calcd.: C, 75.57; H, 6.57; N, 6.30.

Found: C, 75.26; H, 6.68; N, 6.15.

5 Melting point: 206 - 209°C (crystallization solvent :
diisopropyl ether)

Example 99

4-(4-Fluorophenyl)-N-[6-(1-piperidinylmethyl)-7,8-
10 dihydro-2-naphthalenyl]-1-piperidinecarboxamide



6-(1-Pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54 (50 mg, 0.22 mmol) and pyridine (35 mg, 0.44 mmol) were dissolved
15 in tetrahydrofuran (3 ml). Phenyl chlorocarbonate (38 mg, 0.24 mol) was added to the solution under ice-cooling, which was stirred for 10 minutes. The reaction mixture was concentrated, and dimehtylsulfoxide (5 ml) was added to the residue. 4-(4-Fluorophenyl)piperidine hydrochloride (57
20 mg, 0.26 mmol) and 4N aqueous sodium hydroxide solution (0.066 ml, 0.26 mmol) were added to the reaction mixture at room temperature while stirring, which was stirred for 30 minutes. Ethyl acetate and water were added to the mixture, and extraction was conducted. The organic layer
25 was washed with water, and concentrated. Diisopropyl ether was added to the residue. The crystallized product was collected by filtration, washed with diisopropyl ether, to give 4-(4-fluorophenyl)-N-[6-(1-piperidinylmethyl)-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide (48
30 mg) as a white powder.

$^1\text{H-NMR}$ (CDCl_3) δ : 1.60-1.70 (2H, m), 1.79 (4H, m), 1.80-1.90 (2H, m), 2.33 (2H, t, J=7.8 Hz), 2.51 (4H, m), 2.60-2.70 (1H, m), 2.80 (2H, t, J=7.8 Hz), 2.90-3.10 (2H, m), 3.16

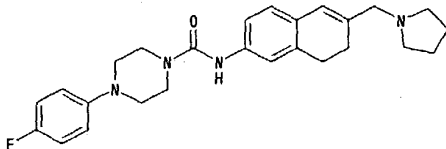
(2H, s), 4.18-4.23 (2H, m), 6.32 (1H, s), 6.32 (1H, s),
6.92-7.09 (4H, m), 7.15-7.20 (3H, m).

Melting point: 182 - 185°C (crystallization solvent :
diisopropyl ether)

5

Example 100

4-(4-Fluorophenyl)-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]-1-piperazinecarboxamide



10 The titled compound was obtained as a white powder in the same manner as in Example 99, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54 and 4-fluorophenylpiperazine.

15 ¹H-NMR (CDCl₃) δ: 1.79 (4H, m), 2.32 (2H, t, J=7.8 Hz), 2.51 (4H, m), 2.80 (2H, t, J=7.8 Hz), 3.13-3.16 (4H, m), 3.16 (2H, s), 3.63-3.66 (4H, m), 6.30 (1H, s), 6.32 (1H, s), 6.88-7.08 (6H, m), 7.19 (1H, s).

Elemental analysis for C₂₆H₃₁FN₄O

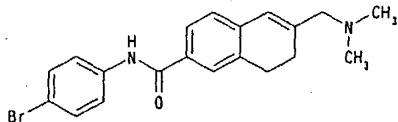
20 Calcd.: C, 71.86; H, 7.19; N, 12.89.

Found: C, 71.68; H, 7.35; N, 12.65.

Melting point: 179 - 181°C (crystallization solvent :
diisopropyl ether)

25 Example 101

N-(4-Bromophenyl)-6-[(dimethylamino)methyl]-7,8-dihydro-2-naphthalenecarboxamide



1) 6-Cyano-1-tetralone (1.30 g, 7.59 mmol)
30 synthesized by a known method by documents (synthetic communications, 23(21), 2965 (1993)) was dissolved in a

mixed solution of concentrated hydrochloric acid (10 ml) and acetic acid (20 ml), which was stirred at 120°C for 16 hours. The reaction mixture was concentrated. Ethyl acetate and water were added to the residue, and extraction was conducted. The organic layer was washed with water, and concentrated. The residue was washed with ethyl acetate - n-hexane (1:1), to give 5-oxo-5,6,7,8-tetrahydro-2-naphthalenecarboxylic acid (1.10 g) as a white powder.

10 ¹H-NMR (CDCl₃) δ: 2.15-2.23 (2H, m), 2.70-2.75 (2H, m), 3.04-3.07 (2H, m), 8.01-8.03 (1H, m), 8.03 (1H, s), 8.13 (1H, d, J=8.7 Hz).

2) N-(4-Bromophenyl)-5-oxo-5,6,7,8-tetrahydro-2-naphthalenecarboxamide (1.21 g) was obtained as a white powder in the same manner as in Example 1, using 5-oxo-5,6,7,8-tetrahydro-2-naphthalenecarboxylic acid (1.00 g, 5.26 mmol) obtained in 1) and 4-bromoaniline (0.90 g, 5.26 mmol).

20 ¹H-NMR (CDCl₃) δ: 2.14-2.23 (2H, m), 2.69-2.73 (2H, m), 3.03-3.07 (2H, m), 7.48-7.58 (4H, m), 7.71 (1H, d, J=8.1 Hz), 7.79 (1H, s), 7.86 (1H, s), 8.12 (1H, d, J=8.1 Hz).

3) N-(4-Bromophenyl)-5-oxo-5,6,7,8-tetrahydro-2-naphthalenecarboxamide (1.10 g, 3.19 mmol) obtained in 2) was dissolved in dimethylformamide diethylacetal (30 ml), which was refluxed with heating for 4 hours. The crystallized product was collected by filtration, washed with ethyl acetate, to give N-(4-bromophenyl)-6-[(dimethylamino)methylidene]-5-oxo-5,6,7,8-tetrahydro-2-naphthalenecarboxamide (1.21 g) as a yellow powder.

30 ¹H-NMR (CDCl₃) δ: 2.80-2.87 (4H, m), 3.07 (6H, m), 7.46-7.72 (7H, m), 7.91 (1H, d, J=8.4 Hz), 8.53 (1H, s).

4) Sodium triacetoxhydroborate (398 mg, 1.87 mmol) was dissolved in a mixed solution of acetic acid (40 ml) and tetrahydrofuran (10 ml) under ice-cooling. N-(4-Bromophenyl)-6-[(dimethylamino)methylidene]-5-oxo-5,6,7,8-tetrahydro-2-naphthalenecarboxamide (500 mg,

35

1.25 mmol) obtained in 3) was added to the solution, which was stirred for 1 hour. The reaction mixture was concentrated under reduced pressure at room temperature.

2-Propanol (50 ml) was added to the residue, and sodium borohydride (142 mg, 3.75 mmol) was further added under ice-cooling. After stirring for 2 hours, the reaction mixture was concentrated. Sodium hydrogencarbonate solution and ethyl acetate was added to the residue for liquid separation. The organic layer was concentrated.

The residue was dissolved in a mixed solution of acetic acid (20 ml) and concentrated hydrochloric acid (20 ml), which was stirred at 70°C for 5 hours. The reaction mixture was concentrated. 4N aqueous sodium hydroxide solution and ethyl acetate were added to the residue, and extraction was conducted. The organic layer was washed with water, and concentrated. The residue was purified by alumina column chromatography (development solvent: ethyl acetate), and the eluent was washed with diisopropyl ether, to give the titled compound (234 mg) as a white powder.

¹H-NMR (CDCl₃) δ: 2.26 (6H, s), 2.38 (2H, t, J=8.1 Hz), 2.89 (2H, t, J=8.1 Hz), 3.02 (2H, s), 6.42 (1H, s), 7.10 (1H, d, J=8.6 Hz), 7.47 (2H, d, J=8.9 Hz), 7.55 (2H, d, J=8.9 Hz), 7.61 (1H, s), 7.62 (1H, d, J=6.7 Hz), 7.76 (1H, s).

Elemental analysis for C₂₀H₂₁BrN₂O

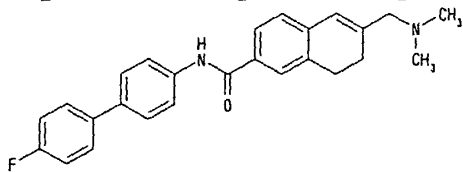
Calcd.: C, 62.35; H, 5.49; N, 7.27.

Found: C, 61.98; H, 5.43; N, 7.07.

Melting point: 175 - 179°C (crystallization solvent : diisopropyl ether)

Example 102

6-[(Dimethylamino)methyl]-N-(4'-fluoro[1,1'-biphenyl]-4-yl)-7,8-dihydro-2-naphthalenecarboxamide



¹H-NMR (CDCl₃) δ: 2.27 (6H, s), 2.39 (2H, t, J=8.4 Hz), 2.91 (2H, t, J=8.4 Hz), 3.02 (2H, s), 6.43 (1H, s), 7.09-7.16 (3H, m), 7.52-7.73 (8H, m), 7.81 (1H, s).

Example 103

Fc1ccc(cc1)-c2ccc(cc2)C(=O)Nc3ccc4ccccc4c3Cc5ccccc5N

¹H-NMR (CDCl₃) δ: 1.75-1.90 (4H, m), 2.36 (2H, t, J=8.1 Hz), 2.45-2.60 (4H, m), 2.85 (2H, t, J=8.1 Hz), 3.18 (2H, s), 6.36 (1H, s), 6.92-7.03 (3H, m), 7.36-7.45 (2H, m), 7.48 (1H, s), 7.62 (2H, d, J=8.4 Hz), 7.78 (1H, s), 7.94 (2H, d, J=8.4 Hz).

Elemental analysis for $C_{28}H_{26}F_2N_2O$

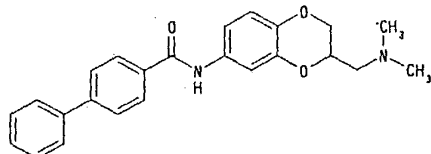
Calcd.: C, 75.66; H, 5.90; N, 6.30.

Found : C, 75.36; H, 5.92; N, 6.10.

Example 104

N-[3-[(Dimethylamino)methyl]-2,3-dihydro-1,4-

benzodioxin-6-yl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained as a white powder by the same method as in Example 1, using N,N-dimethyl-N-[(7-amino-2,3-dihydro-1,4-benzodioxin-2-yl)methyl]amine obtained in Reference Example 62.

¹H-NMR (CDCl₃) δ: 2.33 (6H, s), 2.48-2.66 (2H, m), 3.93-3.99 (1H, m), 4.27-4.31 (2H, m), 6.86 (1H, d, J=8.6 Hz), 7.03-7.07 (1H, m), 7.31-7.32 (1H, m), 7.37-7.49 (3H, m), 7.62 (2H, d, J=7.0 Hz), 7.68 (2H, d, J=8.4 Hz), 7.76 (1H, s), 7.91 (2H, d, J=8.4 Hz).

Elemental analysis for C₂₄H₂₄N₂O₃

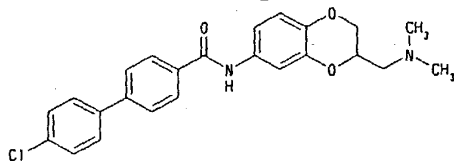
Calcd.: C, 74.21; H, 6.23; N, 7.21.

Found: C, 74.17; H, 6.23; N, 7.01.

Melting point: 124 - 126°C (crystallization solvent : diisopropyl ether)

Example 105

4'-Chloro-N-[3-[(dimethylamino)methyl]-2,3-dihydro-1,4-benzodioxin-6-yl][1,1'-biphenyl]-4-carboxamide



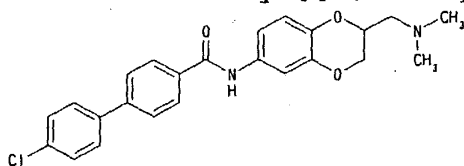
The titled compound was obtained as a white powder by the same method as in Example 1, using N,N-dimethyl-N-[(7-amino-2,3-dihydro-1,4-benzodioxin-2-yl)methyl]amine obtained in Reference Example 62.

¹H-NMR (CDCl₃) δ: 2.33 (6H, s), 2.50-2.67 (2H, m), 3.94-4.01 (1H, m), 4.28-4.31 (2H, m), 6.86 (1H, d, J=8.7 Hz), 7.03-7.06 (1H, m), 7.31 (1H, m), 7.44 (2H, d, J=8.4 Hz), 7.55 (2H, d, J=8.4 Hz), 7.65 (2H, d, J=8.1 Hz), 7.67 (1H, s), 7.91 (2H, d, J=8.1 Hz).

Melting point: 158 - 159°C (crystallization solvent :
diisopropyl ether)

Example 106

- 5 4'-Chloro-N-[2-[(dimethylamino)methyl]-2,3-dihydro-1,4-benzodioxin-6-yl][1,1'-biphenyl]-4-carboxamide



- The titled compound was obtained as a white powder by the same method as in Example 1, using N,N-dimethyl-N-
10 [(6-amino-2,3-dihydro-1,4-benzodioxin-2-yl)methyl]amine obtained in Reference Example 63.

- ¹H-NMR (CDCl₃) δ: 2.34 (6H, s), 2.46-2.67 (2H, m), 3.94-4.01 (1H, m), 4.28-4.34 (2H, m), 6.91 (1H, d, J=8.6 Hz),
7.02-7.05 (1H, m), 7.30 (1H, m), 7.44 (2H, d, J=8.4 Hz),
15 7.55 (2H, d, J=8.4 Hz), 7.66 (2H, d, J=8.1 Hz), 7.70 (1H, s), 7.92 (2H, d, J=8.1 Hz).

Elemental analysis for C₂₄H₂₃ClN₂O₃

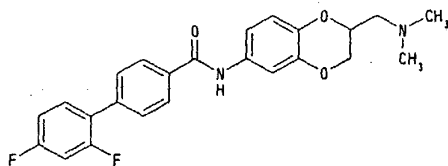
Calcd.: C, 68.16; H, 5.48; N, 6.62.

Found: C, 68.09; H, 5.29; N, 6.57.

- 20 Melting point: 215 - 217°C (crystallization solvent :
diisopropyl ether)

Example 107

- 25 2',4'-Difluoro-N-[2-[(dimethylamino)methyl]-2,3-dihydro-1,4-benzodioxin-6-yl][1,1'-biphenyl]-4-carboxamide



- The titled compound was obtained as a white powder by the same method as in Example 1, using N,N-dimethyl-N-
30 [(6-amino-2,3-dihydro-1,4-benzodioxin-2-yl)methyl]amine

obtained in Reference Example 63.

¹H-NMR (CDCl₃) δ: 2.34 (6H, s), 2.50-2.63 (2H, m), 3.94-4.01 (1H, m), 4.28-4.34 (2H, m), 6.91 (1H, d, J=8.6 Hz), 6.91-7.03 (3H, m), 7.30 (1H, m), 7.40-7.50 (1H, m), 7.61 (2H, d, J=8.1 Hz), 7.69 (1H, s), 7.92 (2H, d, J=8.1 Hz).

Elemental analysis for C₂₄H₂₂F₂N₂O₃

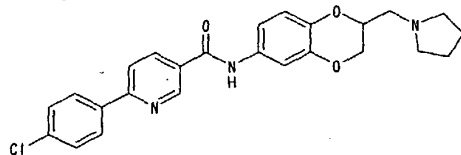
Calcd.: C, 67.91; H, 5.22; N, 6.60.

Found: C, 67.75; H, 5.09; N, 6.48.

Melting point: 209 - 210°C (crystallization solvent : diisopropyl ether)

Example 108

6-(4-Chlorophenyl)-N-[2-(1-pyrrolidinylmethyl)-2,3-dihydro-1,4-benzodioxin-6-yl]nicotinamide



The titled compound was obtained as a white powder by the same method as in Example 1, using 1-[(6-amino-2,3-dihydro-1,4-benzodioxin-2-yl)methyl]pyrrolidine obtained in Reference Example 64.

¹H-NMR (CDCl₃) δ: 1.81 (4H, m), 2.50-2.63 (4H, m), 2.75-2.77 (2H, m), 3.90-4.10 (1H, m), 4.30-4.36 (2H, m), 6.91 (1H, d, J=8.6 Hz), 7.00-7.10 (1H, m), 7.26 (1H, m), 7.48 (2H, d, J=8.6 Hz), 7.72 (1H, s), 7.81 (1H, d, J=7.8 Hz), 8.01 (2H, d, J=8.6 Hz), 8.20-8.25 (1H, m), 9.10 (1H, s).

Elemental analysis for C₂₅H₂₄ClN₃O₃

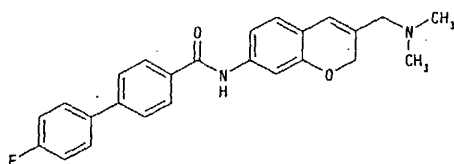
Calcd.: C, 66.74; H, 5.38; N, 9.34.

Found: C, 66.66; H, 5.46; N, 9.11.

Melting point: 218 - 220°C (crystallization solvent : diisopropyl ether)

Example 109

N-[3-[(Dimethylamino)methyl]-2H-chromen-7-yl]-4'-fluoro[1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 3-[(N,N-dimethylamino)methyl]-2H-chromen-7-amine obtained in Reference Example 59.

$^1\text{H-NMR}$ (CDCl_3) δ : 2.23 (6H, s), 2.97 (2H, s), 4.79 (2H, s), 6.30 (1H, s), 6.96 (1H, d, $J=8.1$ Hz), 7.13-7.22 (4H, m), 7.56-7.61 (2H, m), 7.65 (2H, d, $J=8.4$ Hz), 7.78 (1H, s), 7.92 (2H, d, $J=8.4$ Hz).

Elemental analysis for $\text{C}_{25}\text{H}_{23}\text{FN}_2\text{O}_2$

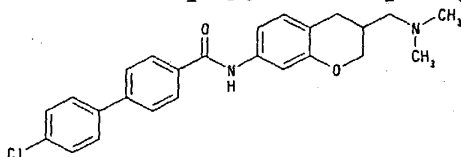
Calcd.: C, 74.61; H, 5.76; N, 6.96.

Found: C, 74.35; H, 5.68; N, 6.74.

Melting point: 192 - 195°C (crystallization solvent : diisopropyl ether)

Example 110

4'-Chloro-N-[3-[(dimethylamino)methyl]-3,4-dihydro-2H-chromen-7-yl][1,1'-biphenyl]-4-carboxamide

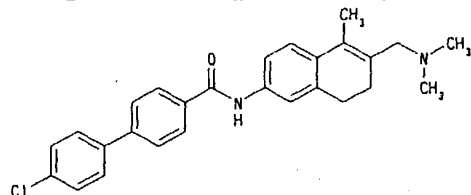


The titled compound was obtained by carrying out the same operation as in Example 1, using N-[(7-amino-3,4-dihydro-2H-chromen-3-yl)methyl]-N,N-dimethylamine obtained in Reference Example 65.

$^1\text{H-NMR}$ (CDCl_3) δ : 2.26 (6H, s), 2.27 (3H, m), 2.47-2.51 (1H, m), 2.83-2.89 (1H, m), 3.82-3.86 (1H, m), 4.28-4.32 (1H, m), 7.04 (1H, d, $J=8.1$ Hz), 7.12-7.18 (2H, m), 7.44 (2H, d, $J=8.4$ Hz), 7.56 (2H, d, $J=8.4$ Hz), 7.67 (2H, d, $J=8.4$ Hz), 7.71 (1H, s), 7.93 (2H, d, $J=8.4$ Hz).

Example 111

4'-Chloro-N-[6-[(dimethylamino)methyl]-5-methyl-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 6-[(dimethylamino)methyl]-5-methyl-7,8-dihydro-2-naphthalenamine obtained in Reference Example 66.

¹H-NMR (CDCl₃) δ: 2.09 (3H, s), 2.27 (6H, s), 2.31-2.37 (2H, m), 2.74-2.79 (2H, m), 3.08 (2H, s), 7.27-7.30 (1H, m), 7.44-7.48 (4H, m), 7.56 (2H, d, J=8.6 Hz), 7.67 (2H, d, J=8.4 Hz), 7.79 (1H, s), 7.95 (2H, d, J=8.4 Hz).

Elemental analysis for C₂₇H₂₇ClN₂O.

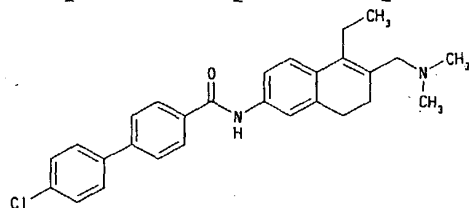
Calcd.: C, 75.25; H, 6.31; N, 6.50.

Found: C, 74.86; H, 6.20; N, 6.42.

Melting point: 199 - 204°C (crystallization solvent : diisopropyl ether)

Example 112

4'-Chloro-N-[6-[(dimethylamino)methyl]-5-ethyl-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 6-[(dimethylamino)methyl]-5-ethyl-7,8-dihydro-2-naphthalenamine obtained in Reference Example 67.

¹H-NMR (CDCl₃) δ: 1.09 (3H, t, J=7.5 Hz), 2.27 (6H, s), 2.31-2.37 (2H, m), 2.60-2.63 (2H, m), 2.71-2.76 (2H, m), 3.08 (2H, s), 7.31 (1H, d, J=9.2 Hz), 7.43-7.49 (4H, m),

7.56 (2H, d, J=8.7 Hz), 7.67 (2H, d, J=8.6 Hz), 7.80 (1H, s), 7.94 (2H, d, J=8.6 Hz).

Elemental analysis for $C_{28}H_{29}ClN_2O$

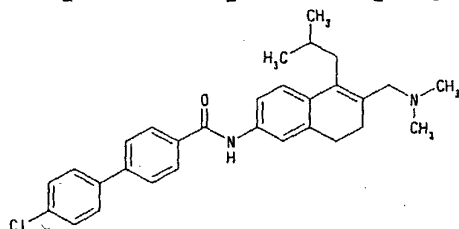
Calcd.: C, 75.57; H, 6.57; N, 6.30.

5 Found: C, 75.41; H, 6.34; N, 6.23.

Melting point: 201 - 204°C (crystallization solvent : diisopropyl ether)

Example 113

10 4'-Chloro-N-[6-[(dimethylamino)methyl]-5-isobutyl-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 6-

15 [(dimethylamino)methyl]-5-isobutyl-7,8-dihydro-2-naphthalenamine obtained in Reference Example 68.

1H -NMR ($CDCl_3$) δ : 0.90 (6H, d, J=6.4 Hz), 1.73-1.78 (1H, m), 2.23 (6H, s), 2.34 (2H, m), 2.50 (2H, d, J=7.3 Hz), 2.74 (2H, m), 3.13 (2H, s), 7.26-7.30 (1H, m), 7.45-7.48
20 (4H, m), 7.56 (2H, d, J=8.7 Hz), 7.67 (2H, d, J=8.4 Hz), 7.79 (1H, s), 7.94 (2H, d, J=8.4 Hz).

Elemental analysis for $C_{30}H_{33}ClN_2O$

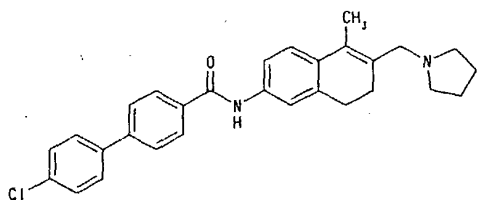
Calcd.: C, 76.17; H, 7.03; N, 5.92.

Found: C, 75.91; H, 7.19; N, 5.72.

25 Melting point: 159 - 162°C (crystallization solvent : diisopropyl ether)

Example 114

30 4'-Chloro-N-[5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



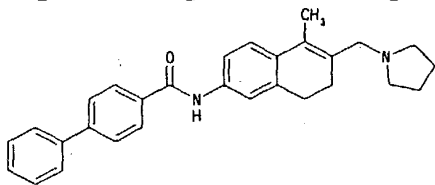
The titled compound was obtained by carrying out the same operation as in Example 1, using 5-methyl-6- (1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine
5 obtained in Reference Example 69.

¹H-NMR (CDCl₃) δ : 1.79 (4H, m), 2.11 (3H, s), 2.30-2.40 (2H, m), 2.54 (4H, m), 2.74-2.79 (2H, m), 3.28 (2H, s), 7.26-7.30 (1H, m), 7.45-7.48 (4H, m), 7.56 (2H, d, J=8.6 Hz), 7.67 (2H, d, J=8.4 Hz), 7.81 (1H, s), 7.95 (2H, d, J=8.4 Hz).
10

Melting point: 190 - 192°C (crystallization solvent : diisopropyl ether)

Example 115

15 N-[5-Methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



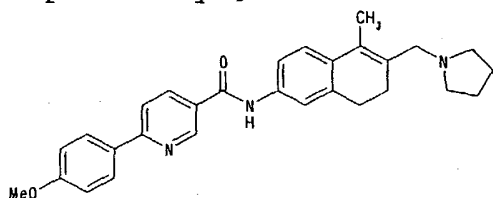
The titled compound was obtained by carrying out the same operation as in Example 1, using 5-methyl-6- (1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine
20 obtained in Reference Example 69.

¹H-NMR (CDCl₃) δ : 1.78 (4H, m), 2.10 (3H, s), 2.35-2.40 (2H, m), 2.53 (4H, m), 2.70-2.78 (2H, m), 3.28 (2H, s), 7.26-7.28 (1H, m), 7.40-7.50 (5H, m), 7.62 (2H, d, J=7.0 Hz), 7.70 (2H, d, J=8.4 Hz), 7.87 (1H, s), 7.94 (2H, d, J=8.4 Hz).
25

Melting point: 169 - 170°C (crystallization solvent : diisopropyl ether)

Example 116

6-(4-Methoxyphenyl)-N-[5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]nicotinamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 69.

¹H-NMR (CDCl₃) δ: 1.78 (4H, m), 2.09 (3H, s), 2.35-2.40 (2H, m), 2.53 (4H, m), 2.70-2.77 (2H, m), 3.27 (2H, s), 3.88 (3H, s), 7.01 (2H, d, J=8.9 Hz), 7.26 (1H, d, J=8.9 Hz), 7.45-7.47 (2H, m), 7.75 (1H, d, J=8.4 Hz), 7.95 (1H, s), 8.01 (2H, d, J=8.9 Hz), 8.18-8.21 (1H, m), 9.09 (1H, m).

Elemental analysis for C₂₉H₃₁N₃O₂

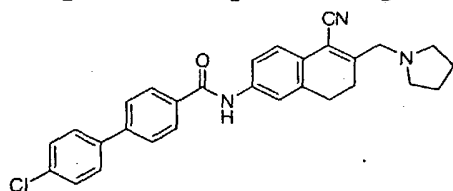
Calcd.: C, 76.79; H, 6.89; N, 9.26.

Found: C, 76.46; H, 6.64; N, 9.09.

Melting point: 165 - 167°C (crystallization solvent : diisopropyl ether)

Example 117

4'-Chloro-N-[5-cyano-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained as a colorless powder by carrying out the same operation as in Example 1, using 6-amino-2-(1-pyrrolidinylmethyl)-3,4-dihydro-1-naphthalenecarbonitrile obtained in Reference Example 70 and 4'-chloro[1,1'-biphenyl]-4-carboxylic acid.

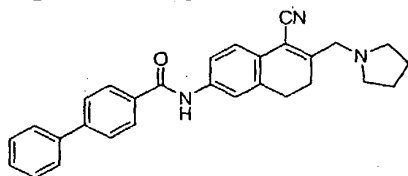
^1H NMR ($\text{DMSO}-d_6$) δ : 1.73 (4H, m), 2.50 (4H, m), 2.56 (2H, m), 2.82 (2H, m), 3.49 (2H, s), 7.32 (1H, d, $J = 9.0$ Hz), 7.57 (2H, d, $J = 8.4$ Hz), 7.56-7.87 (6H, m), 8.07 (2H, d, $J = 8.4$ Hz), 10.40 (1H, s).

5 FABMS(pos) 468.2 $[\text{M}+\text{H}]^+$

Melting point: 191 - 192°C (crystallization solvent : diisopropyl ether)

Example 118

10 N-[5-Cyano-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by as a colorless powder carrying out the same operation as in Example 1, using 6-amino-2-(1-pyrrolidinylmethyl)-3,4-dihydro-1-naphthalenecarbonitrile obtained in Reference Example 70 and [1,1'-biphenyl]-4-carboxylic acid.

^1H NMR ($\text{DMSO}-d_6$) δ : 1.81 (4H, m), 2.62 (6H, m), 2.88 (2H, m), 3.56 (2H, s), 7.41 (2H, m), 7.46 (3H, m), 7.64 (2H, d, $J = 6.9$ Hz), 7.73 (3H, m), 7.88 (1H, s), 7.95 (2H, d, $J = 8.1$ Hz).

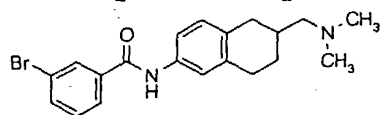
20 FABMS(pos) 434.2 $[\text{M}+\text{H}]^+$

Melting point: 168 - 170°C (crystallization solvent : diisopropyl ether)

25

Example 119

3-Bromo-N-[6-[(dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]benzamide

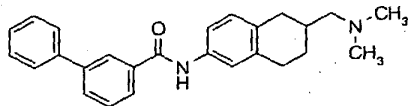


30 The titled compound was obtained by carrying out the same operation as in Example 1, using 6-amino-2-[(N,N-

dimethylamino)methyl]tetralin and 3-bromobenzoic acid.
¹H NMR (DMSO-d₆) δ: 1.31 (1H, m), 1.89 (2H, m), 2.17 (6H, s), 2.17-2.35 (3H, m), 2.77 (3H, m), 7.04 (1H, d, J=8.4 Hz), 7.49 (3H, m), 7.79 (1H, d, J=8.1 Hz), 7.94 (1H, d, J=7.8 Hz), 8.13 (1H, s), 10.20 (1H, s).

Example 120

N-[6-[(Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-3-carboxamide



The titled compound was obtained by carrying out the same operation as in Example 16, using 3-bromo-N-[6-[(dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]benzamide obtained in Example 119 and phenylboronic acid.

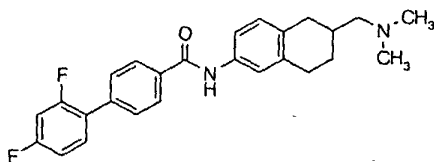
¹H NMR (DMSO-d₆) δ: 1.43 (1H, m), 2.02 (1H, m), 2.21 (1H, m), 2.42 (1H, m), 2.81 (6H, s), 2.88 (3H, m), 3.09 (2H, m), 7.06 (1H, m), 7.42-7.65 (6H, m), 7.78-7.95 (4H, m), 8.22 (1H, s), 10.27 (1H, s).

FABMS(pos) 385.2 [M+H]⁺

Melting point: 145 - 148°C (crystallization solvent : ethyl acetate-diisopropyl ether)

Example 121

N-[6-[(Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-2',4'-difluoro[1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 6-amino-2-[(N,N-dimethylamino)methyl]tetralin and 2',4'-difluoro[1,1'-

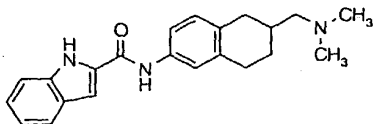
biphenyl]-4-carboxylic acid.

¹H NMR (CDCl₃) δ: 1.41 (1H, m), 1.94 (2H, m), 2.25 (6H, s), 2.23-2.30 (3H, m), 2.86 (3H, m), 6.96 (2H, m), 7.09 (1H, d, J=8.1 Hz), 7.30 (1H, m), 7.43 (2H, m), 7.61 (2H, m), 7.76 (1H, s), 7.93 (2H, m).

Melting point: 162 - 163°C (crystallization solvent : ethyl acetate-diisopropyl ether)

Example 122

10 N-[6-[(Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl-1H-indole-2-carboxamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 6-amino-2-[(N,N-dimethylamino)methyl]tetralin and 1H-indol-2-carboxylic acid.

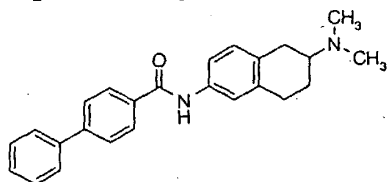
¹H NMR (DMSO-d₆) δ: 1.32 (1H, m), 1.91 (2H, m), 2.16 (6H, s), 2.16-2.35 (3H, m), 2.78 (3H, m), 7.06 (2H, m), 7.21 (1H, m), 7.44 (4H, m), 7.66 (1H, d, J=8.1 Hz), 10.05 (1H, s), 11.68 (1H, s).

FABMS(pos) 348.2 [M+H]⁺

Melting point: 190 - 192°C (crystallization solvent : ethyl acetate - diisopropyl ether)

25 Example 123

N-[6-[(Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl] [1,1'-biphenyl]-4-carboxamide

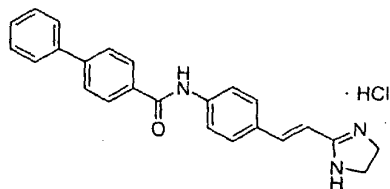


A tetrahydrofuran solution (0.146ml, 0.293mmol) of N-(6-oxo-5,6,7,8-tetrahydro-2-naphthalenyl)[1,1'-

biphenyl]-4-carboxamide (10 mg, 0.029 mmol) obtained in Reference Example 72 and 2N dimethylamine was added to acetic acid-tetrahydrofuran (1:1) solution (0.5ml), which was stirred at 50°C for 15 minutes. After the reaction mixture was cooled at room temperature, sodium triacetoxhydroborate (31 mg, 0.146 mmol) was added, which was stirred at 50°C for 2 hours. 1N Hydrochloric acid was added to the reaction mixture, which was washed with ethyl acetate. Sodium carbonate was added to the water layer to make it alkaline, then extraction was conducted using ethyl acetate. The extract was washed with saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. The resulting residue was purified by alumina B column chromatography (development solvent; ethyl acetate), to give the titled compound (1.6mg).
¹H NMR (CDCl₃) δ: 1.68 (1H, m), 2.27 (1H, m), 2.40 (6H, s), 2.78 (5H, m), 7.11 (1H, d, J=8.1 Hz), 7.32-7.50 (5H, m), 7.62 (2H, m), 7.72 (2H, d, J=8.4 Hz), 7.78 (1H, br), 7.94 (2H, d, J=8.4 Hz).
FABMS(pos) 371.2 [M+H]⁺

Example 124

N-[4-[(E)-2-(4,5-Dihydro-1H-imidazol-2-yl)ethenyl]phenyl][1,1'-biphenyl]-4-carboxamide hydrochloride

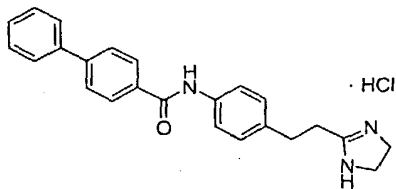


10.1 N Hydrogen chloride-ethanol solution (30 ml) was added to an ethanol suspension of N-[4-[(E)-2-cyanoethenyl]phenyl][1,1'-biphenyl]-4-carboxamide (250 mg, 0.771 mmol) obtained in Reference Example under room temperature, which was stirred for 16 hours. After the

- solvent was distilled out under reduced pressure, ethanol was again added to the residue, and then ethylenediamine (0.155 ml, 2.31 mmol) was added at room temperature, which was stirred for 16 hours. Sodium hydrogencarbonate solution was added to the reaction mixture, and the precipitated crude product was washed with water and chloroform. This product was dissolved in methanol. 1 N Hydrochloric acid (4 ml) was added to the solution, and the solvent was distilled out under reduced pressure.
- Small amount of water was added to the resulting residue, to give the titled compound (124 mg) as a colorless powder. ¹H NMR (DMSO-d₆, free base) δ : 3.33 (4H, m), 6.61 (1H, d, J = 16.8 Hz), 7.15 (1H, d, J = 16.8 Hz), 7.52 (5H, m), 7.83 (6H, m), 8.07 (2H, d, J = 8.4 Hz).
- Elemental analysis for C₂₄H₂₁N₃O · HCl · 1.5H₂O
Calcd.: C, 66.89; H, 5.85; N, 9.75.
Found: C, 67.16; H, 6.10; N, 10.03.

Example 125

- N-[4-[2-(4,5-Dihydro-1H-imidazol-2-yl)ethenyl]phenyl][1,1'-biphenyl]-4-carboxamide hydrochloride



- 10% Palladium - carbon (200 mg) was added to a methanol suspension of N-[4-[(E)-2-(4,5-dihydro-1H-imidazol-2-yl)ethenyl]phenyl][1,1'-biphenyl]-4-carboxamide hydrochloride (80 mg, 0.198 mmol) obtained in Example 124, which was stirred under hydrogen atmosphere at 60°C for 2 hours. After a catalyst was filtered off, the solvent was distilled out under reduced pressure. Diethyl ether was added to the resulting residue, to give the titled compound (52 mg) as a colorless powder.

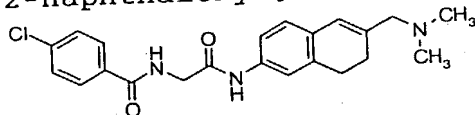
^1H NMR (DMSO- d_6) δ : 2.73-2.97 (4H, m), 3.37 (4H, s), 7.24 (2H, d, $J = 8.4$ Hz), 7.46 (3H, m), 7.76 (6H, m), 8.08 (2H, d, $J = 8.4$ Hz).

FABMS(pos) 370[M+H] $^+$

5

Example 126

4-Chloro-N-[2-[[6-[(dimethylamino)methyl]-7,8-dihydro-2-naphthalenyl]amino]-2-oxoethyl]benzamide



10 The titled compound was obtained by carrying out the same operation as in Example 1, using 6-[(N,N-dimethylamino)methyl]-7,8-dihydro-2-naphthalenamine obtained in Example 41-2) and 4-chlorobenzoyl glycine.

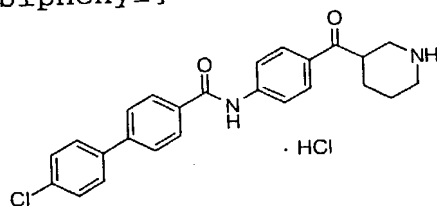
^1H NMR (DMSO- d_6) δ : 2.18 (6H, s), 2.21 (2H, m), 2.71 (2H, m), 2.91 (2H, s), 4.05 (2H, d, $J=5.6$ Hz), 6.30 (1H, s), 6.98 (1H, d, $J=8.1$ Hz), 7.36 (2H, m), 7.58 (2H, d, $J=8.4$ Hz), 7.92 (2H, d, $J=8.4$ Hz), 8.94 (1H, t, $J=5.6$ Hz), 10.00 (1H, s).

FABMS(pos) 398 [M+H] $^+$

20 Melting point: 168 - 171°C (crystallization solvent : diisopropyl ether)

Example 127

25 4'-Chloro-N-[4-(3-piperidinylcarbonyl)phenyl][1,1'-biphenyl]-4-carboxamide hydrochloride



1) tert-Butyl 3-[4-[[4-(4'-chloro[1,1'-biphenyl]-4-yl)carbonyl]amino]benzoyl]-1-piperidinecarboxylate was obtained by carrying out the same operation as in Example 30 1, using tert-butyl 3-(4-aminobenzoyl)-1-piperidinecarboxylate obtained in Reference Example 77 and

4'-chloro[1,1'-biphenyl]-4-carboxylic acid.

FABMS(pos) 519.2 [M+H]⁺

2) 4N Hydrogen chloride-ethyl acetate (1 ml) was added to tert-butyl 3-[4-[[[(4'-chloro[1,1'-biphenyl]-4-yl)carbonyl]amino]benzoyl]-1-piperidinecarboxylate (100 mg, 0.193 mmol) obtained in 1). One hour later, the solvent was distilled out under reduced pressure. Diisopropyl ether was added to the residue, to give the titled compound (73.3 mg) as a colorless powder.

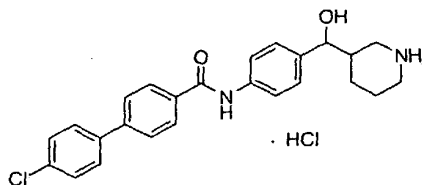
¹H NMR (DMSO-d₆) δ : 1.56 (1H, m), 1.82 (2H, m), 2.02 (1H, m), 2.89 (1H, m), 3.05 (1H, m), 3.33 (2H, m), 3.90 (1H, m), 7.58 (2H, d, J=8.1Hz), 7.81 (2H, d, J=8.1Hz), 7.88 (2H, d, J=8.1Hz), 8.03 (4H, m), 8.11 (2H, d, J=8.1Hz), 9.04 (2H, br), 10.73 (1H, s).

FABMS(pos) 419.2 [M+H]⁺

Melting point: 222 - 225°C (decomposition)

Example 128

4'-Chloro-N-[4-[hydroxy(3-piperidinyl)methyl]phenyl][1,1'-biphenyl]-4-carboxamide hydrochloride



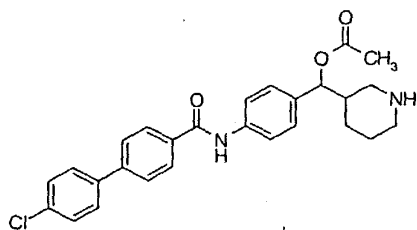
4N Hydrogen chloride-ethyl acetate (1 ml) was added to tert-butyl 3-[[4-[[[(4'-chloro[1,1'-biphenyl]-4-yl)carbonyl]amino]phenyl](hydroxy)methyl]-1-piperidinecarboxylate (100 mg, 0.192 mmol) obtained in Reference Example 78. One hour later, the solvent was distilled out under reduced pressure. Diisopropyl ether was added to the residue, to give the titled compound (79.8 mg) as a colorless powder.

FABMSMS(pos) 421.2 [M+H]⁺

Melting point: 195°C (decomposition)

Example 129

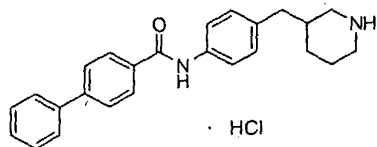
[4-[[[(4'-Chloro[1,1'-biphenyl]-4-yl)carbonyl]amino]phenyl]](3-piperidiny)methyl acetate



5 Concentrated sulfuric acid (0.0562 ml) was added to an acetic acid solution (3.5 ml) of tert-butyl 3-[[4-
10 [[[(4'-chloro[1,1'-biphenyl]-4-yl)carbonyl]amino]phenyl]](hydroxy)methyl]-1-piperidinecarboxylate (366 mg, 0.702 mmol) obtained in Example 128, which was stirred under room temperature for 16 hours. Ethyl acetate was added to the reaction mixture, which was washed with potassium hydrogencarbonate solution and saturated aqueous sodium chloride solution, dried over
15 anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. The resulting oily substance was purified by alumina B column chromatography (development solvent; ethyl acetate : methanol = 3:1), and powdered with diisopropyl ether, to give the titled
20 compound (210 mg).
FABMS(pos) 403.2 [M+H]⁺
Melting point: 200 - 203° C.

Example 130

25 N-[4-(3-Piperidinylmethyl)phenyl][1,1'-biphenyl]-4-carboxamide hydrochloride



4N Hydrogen chloride-ethyl acetate (2 ml) was added to tert-butyl 3-[4-[[[1,1'-biiphenyl]-4-

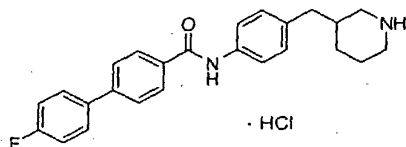
ylcarbonyl)amino]benzyl]-1-piperidinecarboxylate (100 mg, 0.212 mmol) obtained in Reference Example 80. Two hours later, the solvent was distilled out under reduced pressure. Diisopropyl ether was added to the residue for powdering, to give the titled compound (79 mg).

FABMS(pos) 371.3 [M+H]⁺

Melting point: 218 - 220°C (decomposition)

Example 131

4'-Fluoro-N-[4-(3-piperidinylmethyl)phenyl][1,1'-biphenyl]-4-carboxamide hydrochloride



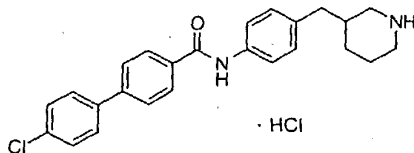
4N Hydrogen chloride-ethyl acetate (3 ml) was added to tert-butyl 3-[4-[[[(4'-fluoro[1,1'-biphenyl]-4-yl)carbonyl]amino]benzyl]-1-piperidinecarboxylate (150 mg, 0.307 mmol) obtained in Reference Example 81. Two hours later, the solvent was distilled out under reduced pressure. Diisopropyl ether was added to the residue, to give the titled compound (115 mg) as a colorless powder.

FABMS(pos) 389.3 [M+H]⁺

Melting point: 205°C (decomposition)

Example 132

4'-Chloro-N-[4-(3-piperidinylmethyl)phenyl][1,1'-biphenyl]-4-carboxamide hydrochloride



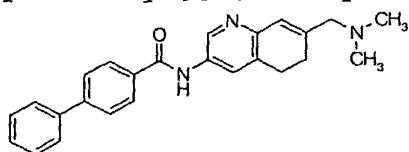
4N Hydrogen chloride-ethyl acetate (3 ml) was added to tert-butyl 3-[4-[[[(4'-chloro[1,1'-biphenyl]-4-yl)carbonyl]amino]benzyl]-1-piperidinecarboxylate (150 mg, 0.297 mmol) obtained in Reference Example 82. Two hours

later, the solvent was distilled out under reduced pressure. Diisopropyl ether was added to the residue, to give the titled compound (73.3 mg) as a colorless powder. FABMS(pos) 405.2 [M+H]⁺

5 Melting point: 200°C (decomposition)

Example 133

N-[7-[(Dimethylamino)methyl]-5,6-dihydro-3-quinolinyl][1,1'-biphenyl]-4-carboxamide



10

The titled compound was obtained by carrying out the same operation as in Example 1, using N-[(3-amino-5,6-dihydro-7-quinolinyl)methyl]-N,N-dimethylamine obtained in Reference Example 86 and [1,1'-biphenyl]-4-carboxylic acid.

15

¹H NMR (DMSO-d₆) δ : 2.16 (6H, s), 2.29 (2H, t, J=8.1 Hz), 2.84 (2H, t, J=8.1 Hz), 2.98 (2H, s), 6.40 (1H, s), 7.42 (1H, m), 7.51 (2H, m), 7.76 (2H, d, J=7.2 Hz), 7.84 (2H, d, J=8.1 Hz), 7.97 (1H, s), 8.06 (2H, d, J=8.4 Hz), 8.65 (1H, s), 10.39 (1H, s).

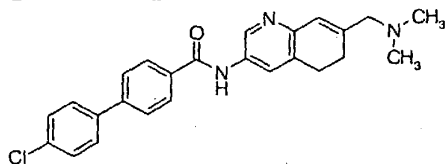
20

FABMS(pos) 384.2 [M+H]⁺

Melting point: 202 - 203°C.

Example 134

25 4'-Chloro-N-[7-[(dimethylamino)methyl]-5,6-dihydro-3-quinolinyl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the same operation as in Example 1, using N-[(3-amino-5,6-dihydro-7-quinolinyl)methyl]-N,N-dimethylamine obtained

30

in Reference Example 86 and 4'-chloro[1,1'-biphenyl]-4-carboxylic acid.

¹H NMR (DMSO-d₆) δ: 2.17 (6H, s), 2.31 (2H, t, J=8.1 Hz), 2.85 (2H, t, J=8.1 Hz), 2.99 (2H, s), 6.41 (1H, s), 7.57 (2H, d, J=8.4 Hz), 7.81 (2H, d, J=8.4 Hz), 7.86 (2H, d, J=8.4 Hz), 7.98 (1H, s), 8.08 (2H, d, J=8.4 Hz), 8.66 (1H, s), 10.41 (1H, s).

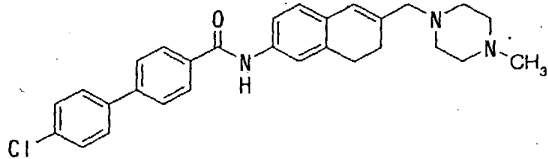
FABMS(pos) 418.2 [M+H]⁺

Melting point: 220 - 222°C.

10

Example 135

4'-Chloro-N-[6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



15

The titled compound was obtained by carrying out the same operation as in Example 51, using 4'-chloro-N-[6-(chloromethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide obtained in Reference Example 56.

¹H-NMR (CDCl₃) δ: 2.30 (3H, s), 2.25-2.50 (10H, m), 2.83 (2H, t, J = 8.1 Hz), 3.07 (2H, s), 6.35 (1H, s), 7.01 (1H, d, J = 8.1 Hz), 7.36 (1H, d, J = 7.8 Hz), 7.44 (2H, d, J = 8.4 Hz), 7.51 (1H, s), 7.55 (2H, d, J = 8.4 Hz), 7.66 (2H, d, J = 8.4 Hz), 7.84 (1H, s), 7.93 (2H, d, J = 8.4 Hz).

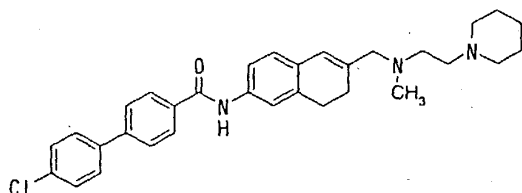
Melting point: 220 - 222°C (crystallization solvent: tetrahydrofuran - n-hexane)

25

Example 136

4'-Chloro-N-[6-[[methyl[2-(1-piperidinyl)ethyl]amino]methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide

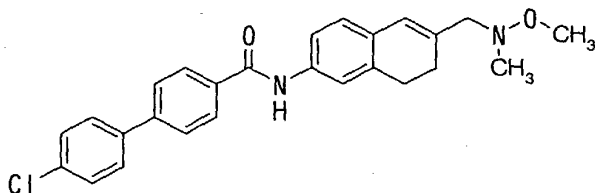
30



5 biphenyl]-4-carboxamide obtained in Reference Example 56.

Melting point: 165 - 167°C (crystallization solvent : tetrahydrofuran - n-hexane)

4'-Chloro-N-[6-[[methoxy(methyl)amino]methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide

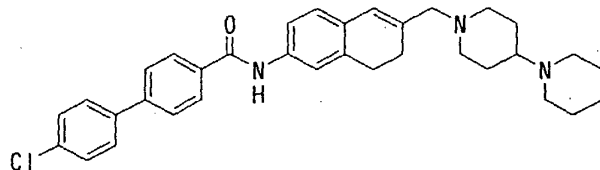


¹H-NMR (CDCl₃) δ : 2.41 (2H, t, J = 8.1 Hz), 2.61 (3H, s),
2.86 (2H, t, J = 8.1 Hz), 3.37 (2H, s), 3.52 (3H, s), 6.39
25 (1H, s), 7.03 (1H, d J = 8.1 Hz), 7.36 (1H, d, J = 8.1 Hz),
7.44 (2H, d, J = 8.4 Hz), 7.53 (1H, s), 7.55 (2H, d, J =
8.4 Hz), 7.66 (2H, d, J = 8.4 Hz), 7.83 (1H, s), 7.93 (2H,
d, J = 8.4 Hz).

Melting point: 190 - 192°C (crystallization solvent :
ethyl acetate - n-hexane)

Example 138

5 4'-Chloro-N-[6-[[4-(1-piperidinyl)-1-piperidinyl]methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide

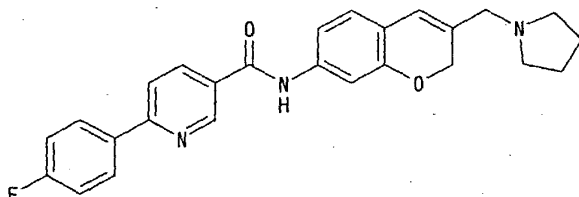


The titled compound was obtained by carrying out the
10 same operation as in Example 51, using 4'-chloro-N-[6-(chloromethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide obtained in Reference Example 56.
¹H-NMR (CDCl₃) δ : 1.45-1.96 (12H, m), 2.29-2.34 (3H, m), 2.57 (4H, s), 2.83 (2H, t, J = 8.1 Hz), 2.96-3.03 (4H, m),
15 6.32 (1H, s), 7.00 (1H, d, J = 8.1 Hz), 7.38 (1H, d, J = 8.1 Hz), 7.44 (2H, d, J = 8.4 Hz), 7.50 (1H, s), 7.55 (2H, d, J = 8.4 Hz), 7.66 (2H, d, J = 8.4 Hz), 7.86 (1H, s), 7.93 (2H, d, J = 8.4 Hz).

Melting point: 232 - 234°C (crystallization solvent :
20 ethyl acetate - n-hexane)

Example 139

6-(4-Fluorophenyl)-N-[3-(1-pyrrolidinylmethyl)-2H-chromen-7-yl]nicotineamide



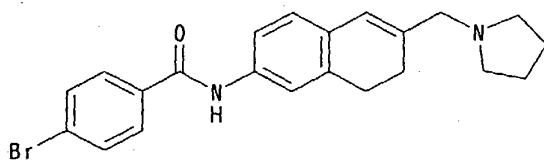
25 The titled compound was obtained by carrying out the same operation as in Example 1, using 3-(1-pyrrolidinylmethyl)-2H-chromen-7-amine obtained in Reference Example 87.

$^1\text{H-NMR}$ (CDCl_3) δ : 1.70 (4H, s), 2.43 (4H, s), 3.12 (2H, s), 4.73 (2H, s), 6.37 (1H, s), 7.03 (1H, d, $J = 7.8$ Hz), 7.29-7.40 (4H, m), 8.15 (1H, d, $J = 8.4$ Hz), 8.22-8.39 (3H, m), 9.15 (1H, s), 10.40 (1H, s).

5 Melting point: 233 - 235°C (crystallization solvent : tetrahydrofuran - n-hexane)

Example 140

10 4-Bromo-N-[6-(1-pyrrolidinymethyl)-7,8-dihydro-2-naphthalenyl]benzamide



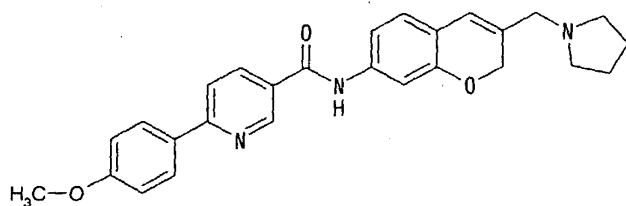
The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(1-pyrrolidinymethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

15 $^1\text{H-NMR}$ (CDCl_3) δ : 1.79 (4H, s), 2.35 (2H, t, $J = 8.1$ Hz), 2.52 (4H, s), 2.83 (2H, t, $J = 8.1$ Hz), 3.17 (2H, s), 6.35 (1H, s), 6.99 (1H, d, $J = 8.1$ Hz), 7.34 (1H, d, $J = 8.1$ Hz), 7.43 (1H, s), 7.60 (2H, d, $J = 8.4$ Hz), 7.72 (2H, d, $J = 8.4$ Hz), 7.76 (1H, s).

20 Melting point: 135 - 137°C (crystallization solvent : ethyl acetate - n-hexane)

Example 141

25 6-(4-Methoxyphenyl)-N-[3-(1-pyrrolidinymethyl)-2H-chromen-7-yl]nicotinamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 3-(1-

pyrrolidinylmethyl)-2H-chromen-7-amine obtained in Reference Example 87.

¹H-NMR (CDCl₃) δ : 1.70 (4H, s), 2.44 (4H, s), 3.12 (2H, s), 3.84 (3H, s), 4.73 (2H, s), 6.37 (1H, s), 7.03 (1H, d, J = 8.1 Hz), 7.09 (2H, t, J = 8.7 Hz), 7.29 (1H, d, J = 8.4 Hz), 7.31 (1H, s), 8.07 (1H, d, J = 8.7 Hz), 8.16 (2H, d, J = 8.7 Hz), 8.32 (1H, d, J = 8.4 Hz), 9.12 (1H, s), 10.34 (1H, s).

Elemental analysis for C₂₇H₂₇N₃O₃

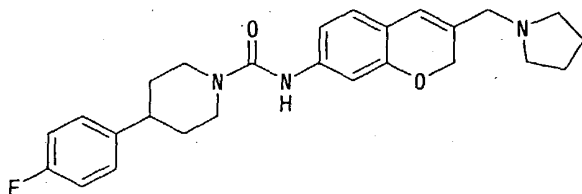
10 Calcd.: C, 73.45; H, 6.16; N, 9.52.

Found: C, 73.02; H, 6.27; N, 9.33.

Melting point: 243 - 245°C (crystallization solvent : tetrahydrofuran - n-hexane)

15 Example 142

4-(4-Fluorophenyl)-N-[3-(1-pyrrolidinylmethyl)-2H-chromen-7-yl]-1-piperidinecarboxamide



20 The titled compound was obtained by carrying out the same operation as in Example 99, using 3-(1-pyrrolidinylmethyl)-2H-chromen-7-amine obtained in Reference Example 87.

¹H-NMR (CDCl₃) δ : 1.69- 1.91 (8H, m), 2.49 (4H, s), 2.70 (1H, t, J = 12.0 Hz), 2.97 (2H, t, J = 12.0 Hz), 3.12 (2H, s), 4.19 (2H, d, J = 13.0 Hz), 4.76 (2H, s), 6.26 (1H, s), 25 6.37 (1H, s), 6.82-7.03 (5H, m), 7.16 (2H, dd, J = 5.4, 8.4 Hz).

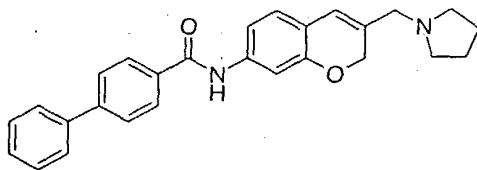
Melting point: 176 - 178°C (crystallization solvent : ethyl acetate - diisopropyl ether)

30

Example 143

N-[3-(1-Pyrrolidinylmethyl)-2H-chromen-7-yl][1,1'-

biphenyl]-4-carboxamide

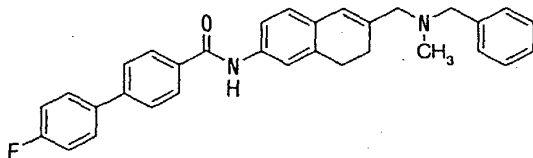


The titled compound was obtained by carrying out the same operation as in Example 1, using 3-(1-pyrrolidinylmethyl)-2H-chromen-7-amine obtained in Reference Example 87.

¹H-NMR (CDCl₃) δ : 1.79 (4H, s), 2.50 (4H, s), 3.15 (2H, s), 4.81 (2H, s), 6.30 (1H, s), 6.95 (1H, d, J = 8.1 Hz), 7.13 (1H, s), 7.20 (1H, d, J = 8.1 Hz), 7.39-7.50 (3H, m), 7.61-7.70 (4H, m), 7.82 (1H, s), 7.92 (2H, d, J = 8.1 Hz).
Melting point: 198 - 200°C (crystallization solvent : ethyl acetate)

Example 144

N-[6-[(N-Benzyl-N-methylamino)methyl]-7,8-dihydro-2-naphthalenyl]-4'-fluoro[1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 6-[(N-benzyl-N-methylamino)methyl]-7,8-dihydro-2-naphthalenamine obtained in Reference Example 88.

¹H-NMR (CDCl₃) δ : 2.20 (3H, s), 2.38 (2H, t, J = 8.1 Hz), 2.85 (2H, t, J = 8.1 Hz), 3.09 (2H, s), 3.52 (2H, s), 6.39 (1H, s), 7.02 (1H, d, J = 8.1 Hz), 7.13-7.66 (13H, m), 7.84 (1H, s), 7.93 (2H, d, J = 8.4 Hz).
Melting point: 143 - 145°C (crystallization solvent : ethyl acetate - n-hexane)

Example 145

4'-Isobutyrylamino-N-[6-(1-pyrrolidinylmethyl)-7,8-

CC(C)C(=O)Nc1ccc(cc1)-c2ccc(cc2)C(=O)Nc3ccc4ccccc4c3CN5CCCC5

Example 146

CCOC(=O)c1ccc(cc1)-c2ccc(cc2)C(=O)Nc3ccc4ccccc4c3CN5CCCCC5

20 Example 147

OC(=O)CCc1ccc(cc1)-c2ccc(cc2)C(=O)Nc3ccc4ccccc4c3CN5CCCC5

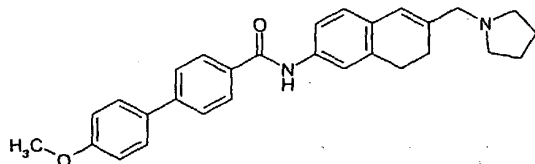
25 The titled compound was obtained as a powder by carrying out the same operation as in Example 1, using

6-(1-pyrrolidinylmethyl)-7,8--dihydro-2-naphthalenamine
obtained in Reference Example 54.

MS m/z 481.4 (MH^+).

5 Example 148

4'-Methoxy-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



10 The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

1H -NMR ($CDCl_3$) δ : 1.80 (4H, m), 2.36 (2H, t, $J=7.8$ Hz), 2.52 (4H, m), 2.86 (2H, t, $J=7.8$ Hz), 3.18 (2H, s), 3.87 (3H, s), 6.36 (1H, s), 7.00-7.03 (3H, m), 7.26 (1H, m), 7.38 (1H, d, $J=8.3$ Hz), 7.49 (1H, s), 7.58 (2H, d, $J=8.6$ Hz), 7.67 (1H, d, $J=8.2$ Hz), 7.78 (1H, s), 7.90 (2H, d, $J=8.2$ Hz).

Elemental analysis for $C_{29}H_{30}N_2O_2$

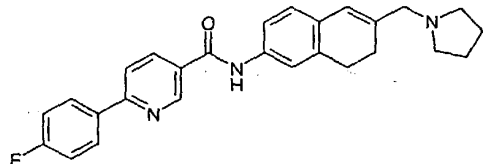
Calcd.: C, 79.42; H, 6.89; N, 6.39.

20 Found: C, 79.21; H, 6.88; N, 6.35.

Melting point: 187-188 $^{\circ}C$ (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 149

25 6-(4-Fluorophenyl)-N-[6-[(1-pyrrolidinyl)methyl]-7,8-dihydro-2-naphthalenyl]nicotinamide



30 The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine

obtained in Reference Example 54.

¹H-NMR (CDCl₃) δ: 1.81 (4H, m), 2.36 (2H, t, J=8.1 Hz), 2.53 (4H, m), 2.86 (2H, t, J=8.1 Hz), 3.18 (2H, s), 6.37 (1H, s), 7.03 (1H, d, J=7.8 Hz), 7.16-7.30 (3H, m), 7.47 (1H, s), 7.77-7.82 (2H, m), 8.06 (2H, dd, J=8.9, 5.3 Hz), 8.25 (1H, dd, J=8.4, 2.2 Hz), 9.11 (1H, d, J=2.0 Hz).

Elemental analysis for C₂₇H₂₆FN₃O

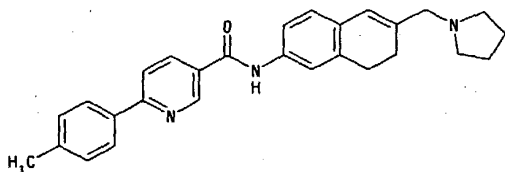
Calcd.: C, 75.85; H, 6.13; N, 9.83.

Found: C, 75.71; H, 5.93; N, 9.75.

Melting point: 225-227 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 150

6-(4-Methylphenyl)-N-[6-[(1-pyrrolidinyl)methyl]-7,8-dihydro-2-naphthalenyl]nicotinamide



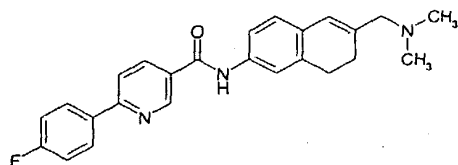
The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

¹H-NMR (CDCl₃) δ: 1.81 (4H, m), 2.36 (2H, t, J=7.8 Hz), 2.43 (3H, s), 2.53 (4H, m), 2.86 (2H, t, J=7.8 Hz), 3.19 (2H, s), 6.37 (1H, s), 7.02 (1H, d, J=8.7 Hz), 7.25-7.39 (3H, m), 7.47 (1H, s), 7.82 (2H, m), 7.96 (2H, d, J=8.1 Hz), 8.23 (1H, dd, J=8.1, 2.3 Hz), 9.12 (1H, d, J=2.3 Hz).

Melting point: 235-236 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 151

N-[6-[(Dimethylamino)methyl]-7,8-dihydro-2-naphthalenyl]-6-(4-fluorophenoxy)nicotinamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 6-[(N,N-dimethylamino)methyl]-7,8-dihydro-2-naphthalenamine
 5 obtained in Reference Example 41-2).

¹H-NMR (CDCl₃) δ: 2.25 (6H, s), 2.34 (2H, t, J=8.1 Hz), 2.86 (2H, t, J=8.1 Hz), 2.99 (2H, s), 6.35 (1H, s), 7.03 (1H, d, J=8.1 Hz), 7.17 (2H, m), 7.26 (1H, m), 7.39 (1H, d, J=8.1 Hz), 7.47 (1H, s), 7.78 (1H, d, J=7.2 Hz), 7.83
 10 (1H, s), 8.06 (1H, dd, J=8.4, 6.7 Hz), 8.25 (1H, d, J=6.7 Hz), 9.12 (1H, s).

Elemental analysis for C₂₅H₂₄FN₃O

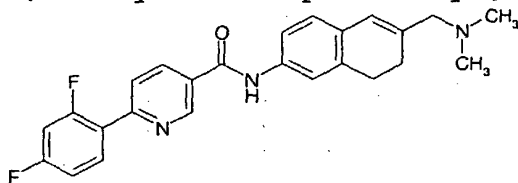
Calcd.: C, 74.79; H, 6.03; N, 10.47.

Found: C, 74.74; H, 5.95; N, 10.24.

15 Melting point: 216-219 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 152

6-(2,4-Difluorophenyl)-N-[6-[(dimethylamino)methyl]-7,8-dihydro-2-naphthalenyl]nicotinamide
 20



The titled compound was obtained by carrying out the same operation as in Example 1, using 6-[(N,N-dimethylamino)methyl]-7,8-dihydro-2-naphthalenamine
 25 obtained in Reference Example 41-2).

¹H-NMR (CDCl₃) δ: 2.25 (6H, s), 2.34 (2H, t, J=8.1 Hz), 2.85 (2H, t, J=8.1 Hz), 3.00 (2H, s), 6.35 (1H, s), 6.90-7.06 (3H, m), 7.39 (1H, d, J=7.8 Hz), 7.47 (1H, s), 7.80-7.90 (2H, m), 8.10 (1H, dd, J=15.3, 8.8 Hz), 8.23 (1H, dd, J=8.4,
 30 2.3 Hz), 9.15 (1H, d, J=1.7 Hz).

Elemental analysis for $C_{25}H_{23}F_2N_3O$

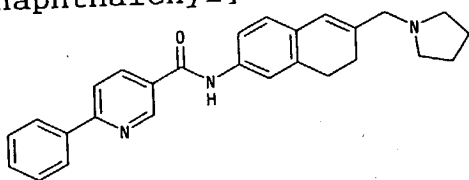
Calcd.: C, 71.58; H, 5.53; N, 10.02.

Found: C, 71.50; H, 5.49; N, 9.61.

Melting point: 162-163 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 153

6-Phenyl-N-[6-(1-pyrrolidinymethyl)-7,8-dihydro-2-naphthalenyl]nicotinamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(1-pyrrolidinymethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

1H -NMR ($CDCl_3$) δ : 1.81 (4H, m), 2.36 (2H, t, $J=8.1$ Hz), 2.53 (4H, m), 2.85 (2H, t, $J=8.1$ Hz), 3.18 (2H, s), 6.37 (1H, s), 7.02 (1H, d, $J=8.1$ Hz), 7.37-7.53 (5H, m), 7.83 (1H, d, $J=8.1$ Hz), 7.86 (1H, d, $J=6.2$ Hz), 8.04 (1H, s), 8.06 (1H, d, $J=1.7$ Hz), 8.24 (1H, dd, $J=8.4, 2.4$ Hz), 9.13 (1H, d, $J=2.2$ Hz).

Elemental analysis for $C_{27}H_{27}N_3O$

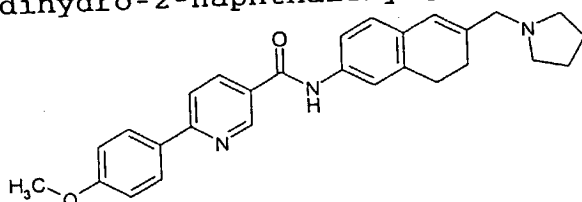
Calcd.: C, 79.19; H, 6.65; N, 10.26.

Found: C, 78.93; H, 6.65; N, 10.19.

Melting point: 186-187 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 154

6-(4-Methoxyphenyl)-N-[6-(1-pyrrolidinymethyl)-7,8-dihydro-2-naphthalenyl]nicotinamide

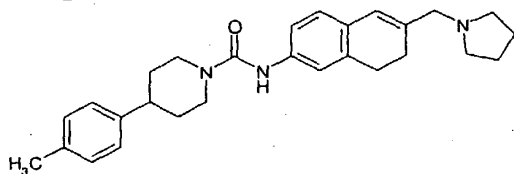


The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

- 5 ¹H-NMR (CDCl₃) δ: 1.80 (4H, m), 2.36 (2H, t, J=8.1 Hz), 2.52 (4H, m), 2.84 (2H, t, J=8.1 Hz), 3.18 (2H, s), 3.88 (3H, s), 6.36 (1H, s), 7.02 (3H, m), 7.37 (1H, d, J=7.5 Hz), 7.47 (1H, s), 7.78 (1H, d, J=8.1 Hz), 7.79 (1H, s), 8.03 (2H, d, J=8.5 Hz), 8.20 (1H, d, J=8.1 Hz), 9.08 (1H, s).
- 10 Melting point: 219-220 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 155

- 4-(4-Methylphenyl)-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide



The titled compound was obtained by carrying out the same operation as in Example 99, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

- 20 ¹H-NMR (CDCl₃) δ: 1.64-1.92 (8H, m), 2.29 (2H, m), 2.32 (3H, s), 2.51 (4H, m), 2.64 (1H, m), 2.80 (2H, t, J=7.8 Hz), 2.97 (2H, dd, J=13.1, 10.7 Hz), 3.15 (2H, s), 4.19 (2H, d, J=13.1 Hz), 6.32 (1H, s), 6.35 (1H, s), 6.42 (1H, d, J=7.8 Hz), 7.06-7.20 (6H, m)

Elemental analysis for C₂₈H₃₅N₃O · 0.5H₂O

Calcd.: C, 76.67; H, 8.27; N, 9.58.

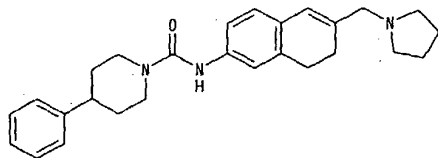
Found: C, 76.72; H, 8.03; N, 9.36.

- 30 Melting point: 197-198 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 156

4-Phenyl-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-

naphthalenyl]-1-piperidinecarboxamide



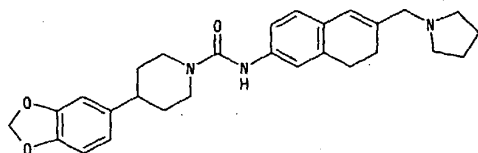
The titled compound was obtained by carrying out the same operation as in Example 99, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

¹H-NMR (CDCl₃) δ: 1.72-1.94 (8H, m), 2.32 (2H, t, J=8.1 Hz), 2.50 (4H, m), 2.72 (1H, m), 2.80 (2H, t, J=8.1 Hz), 2.99 (2H, dd, J=13.4, 10.6 Hz), 3.16 (2H, s), 4.21 (2H, d, J=13.4 Hz), 6.32 (1H, s), 6.34 (1H, s), 6.93 (1H, d, J=8.4 Hz), 7.07 (1H, d, J=8.1 Hz), 7.20-7.35 (6H, m).

Melting point: 184-186 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 157

4-(1,3-Benzodioxol-5-yl)-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide



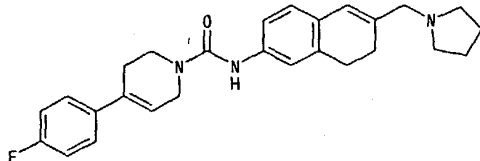
The titled compound was obtained by carrying out the same operation as in Example 99, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

¹H-NMR (CDCl₃) δ: 1.61-1.88 (8H, m), 2.31 (2H, t, J=8.1 Hz), 2.51 (4H, m), 2.59 (1H, m), 2.62 (2H, t, J=8.1 Hz), 2.94 (2H, dd, J=13.1, 11.2 Hz), 3.15 (2H, s), 4.18 (2H, d, J=13.1 Hz), 5.93 (2H, s), 6.31 (1H, s), 6.44 (1H, s), 6.64-6.77 (3H, m), 6.92 (1H, d, J=8.1 Hz), 7.07 (1H, d, J=8.1 Hz), 7.19 (1H, s).

Melting point: 149-150 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 158

4-(4-Fluorophenyl)-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]-3,6-dihydro-1(2H)-pyridinecarboxamide



The titled compound was obtained by carrying out the same operation as in Example 99, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

$^1\text{H-NMR}$ (CDCl_3) δ : 1.79 (4H, m), 2.32 (2H, t, $J=8.1$ Hz), 2.50 (4H, m), 2.59 (2H, brt), 2.80 (2H, t, $J=8.1$ Hz), 3.17 (2H, s), 3.74 (2H, t, $J=5.7$ Hz), 4.15 (2H, d, $J=2.5$ Hz), 6.00 (1H, brt), 6.32 (1H, s), 6.32 (1H, s), 6.94 (1H, d, $J=8.1$ Hz), 7.00-7.32 (6H, m).

Elemental analysis for $\text{C}_{27}\text{H}_{30}\text{FN}_3\text{O}$

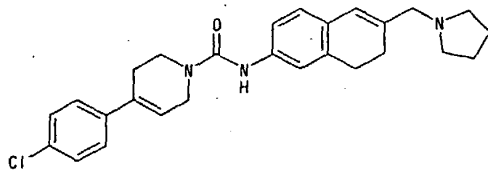
Calcd.: C, 75.15; H, 7.01; N, 9.74.

Found: C, 75.09; H, 6.93; N, 9.77.

Melting point: 206-207 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 159

4-(4-Chlorophenyl)-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]-3,6-dihydro-1(2H)-pyridinecarboxamide



The titled compound was obtained by carrying out the same operation as in Example 99, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

¹H-NMR (CDCl₃) δ: 1.79 (4H, m), 2.32 (2H, t, J=8.1 Hz), 2.50 (4H, m), 2.59 (2H, brt), 2.80 (2H, t, J=8.1 Hz), 3.16 (2H, s), 3.73 (2H, t, J=5.6 Hz), 4.15 (2H, d, J=2.8 Hz), 6.06 (1H, brt), 6.30 (1H, s), 6.32 (1H, s), 6.93 (1H, d, J=7.8 Hz), 7.09 (1H, d, J=7.8 Hz), 7.21-7.31 (5H, m).

Elemental analysis for C₂₇H₃₀ClN₃O

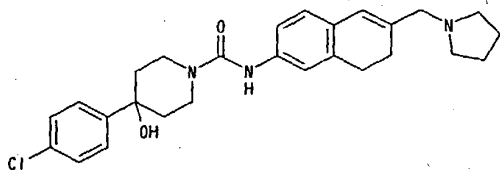
Calcd.: C, 72.39; H, 6.75; N, 9.38.

Found: C, 72.19; H, 6.75; N, 9.19.

Meltingpoint: 217-218 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 160

4-(4-Chlorophenyl)-4-hydroxy-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide



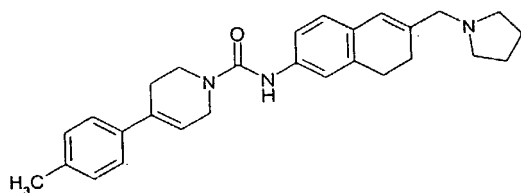
The titled compound was obtained by carrying out the same operation as in Example 99, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

¹H-NMR (CDCl₃) δ: 1.79 (4H, m), 1.80 (2H, m), 2.04 (1H, dd, J=13.1, 10.8 Hz), 2.06 (1H, dd, J=13.1, 10.8 Hz), 2.31 (2H, t, J=7.8 Hz), 2.50 (1H, brs), 2.51 (4H, m), 2.79 (2H, t, J=7.8 Hz), 3.15 (2H, s), 3.41 (2H, dd, J=12.6, 10.8 Hz), 4.00 (2H, d, J=12.6 Hz), 6.32 (1H, s), 6.37 (1H, s), 6.93 (1H, d, J=8.1 Hz), 7.05-7.42 (6H, m).

Meltingpoint: 181-182 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 161

4-(4-Methylphenyl)-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]-3,6-dihydro-1(2H)-pyridinecarboxamide



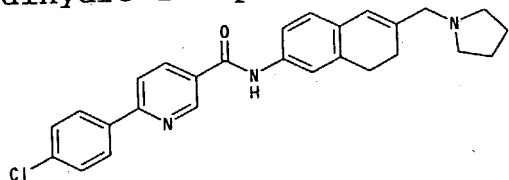
The titled compound was obtained by carrying out the same operation as in Example 99, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

¹H-NMR (CDCl₃) δ: 1.79 (4H, m), 2.32 (2H, t, J=7.8 Hz), 2.35 (3H, s), 2.50 (4H, m), 2.61 (2H, brt), 2.80 (2H, t, J=7.8 Hz), 3.16 (2H, s), 3.73 (2H, t, J=5.7 Hz), 4.15 (2H, d, J=2.8 Hz), 6.03 (1H, s), 6.29 (1H, s), 6.32 (1H, s), 6.93 (1H, d, J=8.1 Hz), 7.07-7.30 (6H, m).

Melting point: 199-202 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 162

6-(4-Chlorophenyl)-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]nicotinamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

¹H-NMR (CDCl₃+DMSO-d₆) δ: 1.80 (4H, m), 2.32-2.58 (6H, m), 2.85 (2H, t, J=8.0 Hz), 3.18 (2H, s), 6.36 (1H, s), 7.01 (1H, d, J=8.4 Hz), 7.48 (2H, d, J=8.4 Hz), 7.49 (1H, m), 7.59 (1H, s), 7.83 (1H, d, J=8.4 Hz), 8.04 (2H, d, J=8.4 Hz), 8.35 (1H, dd, J=8.4, 2.2 Hz), 9.25 (1H, d, J=2.2 Hz), 9.42 (1H, s).

Elemental analysis for C₂₇H₂₆ClN₃O

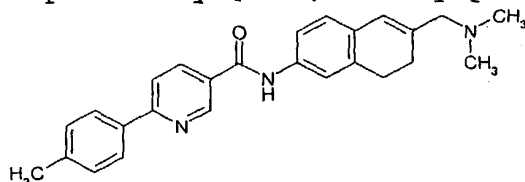
Calcd.: C, 73.04; H, 5.90; N, 9.46.

Found: C, 73.11; H, 5.71; N, 9.20.

Melting point: 252-253 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 163

- 5 N-[6-[(Dimethylamino)methyl]-7,8-dihydro-2-naphthalenyl]-6-(4-methylphenyl)nicotinamide



- 10 The titled compound was obtained by carrying out the same operation as in Example 1, using 6-[(N,N-dimethylamino)methyl]-7,8-dihydro-2-naphthalenamine obtained in Reference Example 41-2).

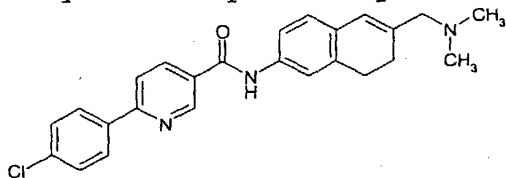
- ¹H-NMR (CDCl₃) δ: 2.25 (6H, s), 2.34 (2H, t, J=8.1 Hz), 2.43 (3H, s), 2.85 (2H, t, J=8.1 Hz), 2.99 (2H, s), 6.34 (1H, s), 7.02 (1H, d, J=8.1 Hz), 7.31 (2H, d, J=8.1 Hz), 7.39 (1H, d, J=8.1 Hz), 7.46 (1H, s), 7.81 (1H, d, J=8.4 Hz), 7.87 (1H, s), 7.96 (2H, d, J=8.1 Hz), 8.22 (1H, dd, J=8.4, 2.3 Hz), 9.11 (1H, d, J=2.3 Hz).

- 15 Melting point: 228-230 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

20

Example 164

- 6-(4-Chlorophenyl)-N-[6-[(dimethylamino)methyl]-7,8-dihydro-2-naphthalenyl]nicotinamide



- 25 The titled compound was obtained by carrying out the same operation as in Example 1, using 6-[(N,N-dimethylamino)methyl]-7,8-dihydro-2-naphthalenamine obtained in Reference Example 41-2).

- ¹H-NMR (CDCl₃) δ: 2.25 (6H, s), 2.35 (2H, t, J=8.1 Hz), 2.86 (2H, t, J=8.1 Hz), 2.99 (2H, s), 6.35 (1H, s), 7.04 (1H,
- 30

d, J=8.1 Hz), 7.40 (1H, d, J=8.4 Hz), 7.49 (1H, brs), 7.49 (2H, d, J=8.4 Hz), 7.78 (1H, s), 7.84 (1H, d, J=8.4 Hz), 8.02 (2H, d, J=8.4 Hz), 8.26 (1H, dd, J=8.1, 2.2 Hz), 9.13 (1H, d, J=2.2 Hz).

5 Elemental analysis for $C_{25}H_{24}ClN_3O$

Calcd.: C, 71.85; H, 5.79; N, 10.05.

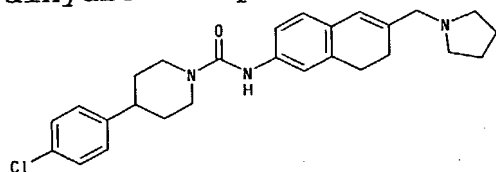
Found: C, 71.88; H, 5.67; N, 9.86.

Melting point: 248-249 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

10

Example 165

4-(4-Chlorophenyl)-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide



15

The titled compound was obtained by carrying out the same operation as in Example 99, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

1H -NMR ($CDCl_3$) δ : 1.66-1.91 (8H, m), 2.32 (2H, t, J=8.1 Hz), 2.50 (4H, m), 2.70 (1H, m), 2.80 (2H, t, J=8.1 Hz), 2.98 (2H, dd, J=13.7, 12.0 Hz), 3.16 (2H, s), 4.20 (2H, d, J=13.7 Hz), 6.32 (1H, s), 6.32 (1H, s), 6.93 (1H, d, J=8.1 Hz), 7.05-7.30 (6H, m).

Elemental analysis for $C_{27}H_{32}ClN_3O$

25 Calcd.: C, 72.06; H, 7.17; N, 9.34.

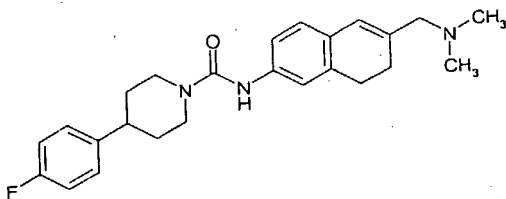
Found: C, 72.08; H, 7.23; N, 9.15.

Melting point: 194-195 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

30

Example 166

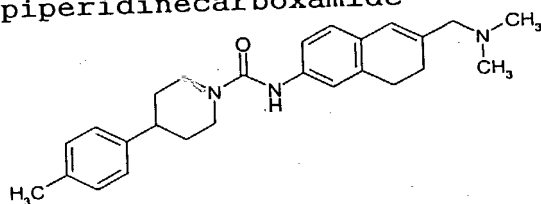
N-[6-[(Dimethylamino)methyl]-7,8-dihydro-2-naphthalenyl]-4-(4-fluorophenyl)-1-piperidinecarboxamide



- The titled compound was obtained by carrying out the same operation as in Example 99, using 6-[(N,N-dimethylamino)methyl]-7,8-dihydro-2-naphthalenamine
- 5 obtained in Reference Example 41-2).
- ¹H-NMR (CDCl₃) δ: 1.65-1.75 (2H, m), 1.89 (2H, d, J=11.4 Hz), 2.23 (6H, s), 2.30 (2H, t, J=8.1 Hz), 2.70 (1H, m), 2.80 (2H, t, J=8.1 Hz), 2.94-3.01 (4H, m), 4.20 (2H, d, J=13.4 Hz), 6.30 (1H, s), 6.35 (1H, s), 6.92-7.20 (7H, m).
- 10 Melting point: 187-188 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 167

- N-[6-[(Dimethylamino)methyl]-7,8-dihydro-2-naphthalenyl]-4-(4-methylphenyl)-1-piperidinecarboxamide
- 15

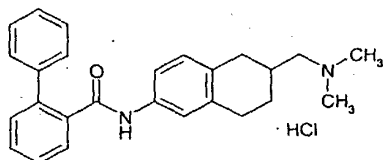


- The titled compound was obtained by carrying out the same operation as in Example 99, using 6-[(N,N-dimethylamino)methyl]-7,8-dihydro-2-naphthalenamine
- 20 obtained in Reference Example 41-2).
- ¹H-NMR (CDCl₃) δ: 1.66-1.74 (2H, m), 1.89 (2H, d, J=11.7 Hz), 2.28 (6H, s), 2.30 (2H, t, J=8.1 Hz), 2.38 (3H, s), 2.68 (1H, m), 2.80 (2H, t, J=8.1 Hz), 2.94-3.02 (4H, m), 4.19 (2H, d, J=12.8 Hz), 6.30 (1H, s), 6.35 (1H, s), 6.93
- 25 (1H, d, J=8.1 Hz), 7.07-7.20 (6H, m).
- Elemental analysis for C₂₆H₃₃N₃O · 0.5H₂O
- Calcd.: C, 75.69; H, 8.31; N, 10.18
- Found: C, 75.44; H, 8.16; N, 10.05
- 30 Melting point: 200-202 °C (crystallization solvent: ethyl

acetate - diisopropyl ether)

Example 168

5 N-[6-[(Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-2-carboxamide hydrochloride



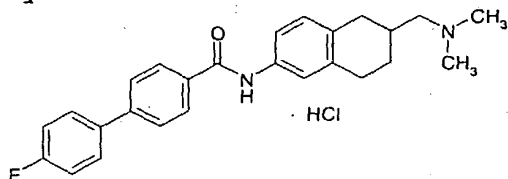
The titled compound was obtained by carrying out the same operation as in Example 1, using 6-amino-2-[(N,N-dimethylamino)methyl]tetralin hydrochloride.

10 ¹H-NMR (DMSO-d₆) δ: 1.39 (1H, m), 1.99 (1H, m), 2.17 (1H, m), 2.42 (1H, dd, J=16.2, 10.1 Hz), 2.78 (6H, s), 2.88 (1H, dd, J=16.2, 4.5 Hz), 3.06 (2H, t, J=5.7 Hz), 3.38 (2H, s), 6.94-7.62 (11H, m), 7.64 (1H, d, J=1.7 Hz), 10.11 (1H, brs), 10.18 (1H, s).

15 Melting point: 196-197 °C (crystallization solvent: methanol - ethyl acetate)

Example 169

20 N-[6-[(Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-4'-fluoro[1,1'-biphenyl]-4-carboxamide hydrochloride



25 4'-Fluoro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide synthesized in Example 42 was dissolved in ethyl acetate. An excess amount of 4N hydrochloric acid-ethyl acetate solution was added to the solution, which was concentrated under reduced pressure. The resulting residue was recrystallized from methanol - ethyl

30

acetate, to give the titled compound.

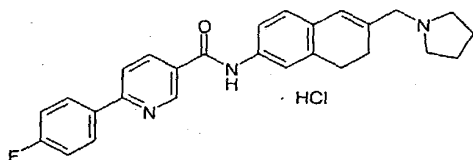
¹H-NMR (DMSO-d₆) δ: 1.43 (1H, m), 2.06 (1H, m), 2.21 (1H, m), 2.45 (1H, m), 2.79 (6H, s), 2.92 (1H, dd, J=16.2, 4.2 Hz), 3.08 (2H, d, J=6.4 Hz), 3.33 (2H, s), 7.05 (1H, d, J=8.4 Hz), 7.34 (2H, dd, J=8.9, 8.9 Hz), 7.53 (1H, d, J=8.4 Hz), 7.59 (1H, s), 7.80 (4H, m), 8.06 (2H, d, J=8.1 Hz), 10.02 (1H, s), 10.03 (1H, brs).

Melting point:: 240-245 °C (crystallization solvent: methanol - ethyl acetate)

10

Example 170

6-(4-Fluorophenyl)-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]nicotinamide hydrochloride



15 The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

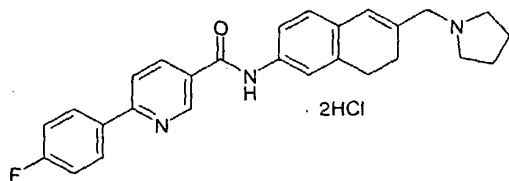
¹H-NMR (DMSO-d₆) δ: 1.70 (4H, m), 2.26 (2H, t, J=8.1 Hz), 2.44 (4H, m), 2.76 (2H, t, J=8.1 Hz), 3.12 (2H, s), 3.34 (1H, s), 6.36 (1H, s), 7.03 (1H, d, J=7.8 Hz), 7.37 (2H, dd, J=8.4, 7.0 Hz), 7.57 (1H, d, J=8.4 Hz), 7.59 (1H, s), 8.13-8.42 (4H, m), 9.19 (1H, s), 10.43 (1H, s).

20 Melting point: 229-231 °C (crystallization solvent: methanol - ethyl acetate)

25

Example 171

6-(4-Fluorophenyl)-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]nicotinamide dihydrochloride



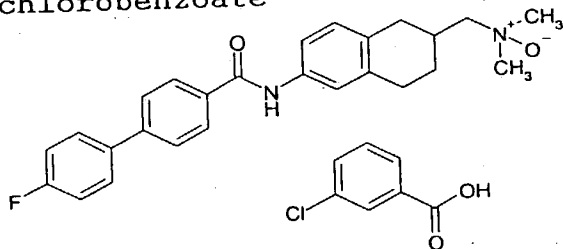
30

The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

- 5 ¹H-NMR (DMSO-d₆) δ: 2.00 (4H, m), 2.45 (4H, m), 2.83 (2H, t, J=8.1 Hz), 3.05 (2H, m), 3.47 (2H, m), 3.88 (1H, s), 6.69 (1H, s), 7.13 (1H, d, J=8.1 Hz), 7.38 (2H, dd, J=8.9, 8.6 Hz), 7.64 (1H, d, J=10.6 Hz), 7.66 (1H, s), 8.14-8.42 (4H, m), 9.19 (1H, s), 10.52 (1H, s), 10.60 (1H, brs).
- 10 Melting point: 245-248 °C (crystallization solvent: methanol - ethyl acetate)

Example 172

- 15 N-[6-[(Dimethylnitroyl)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-4'-fluoro[1,1'-biphenyl]-4-carboxamide 3-chlorobenzoate

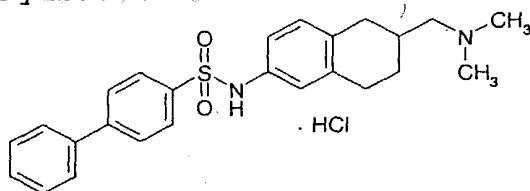


- 4'-FluoroN-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide (100 mg) obtained in Example 42 was dissolved in acetone (10 ml), which was stirred under ice-cooling. 3-Chloroperbenzoic acid (purity : 50%) (86 mg) was added to the solution, which was stirred under ice-cooling for 1 hour. The reaction mixture was concentrated under reduced pressure, and the residue was washed with diisopropyl ether, to give the titled compound (158 mg).
- 20 ¹H-NMR (DMSO-d₆) δ: 1.57 (1H, m), 2.07 (1H, m), 2.61 (1H, m), 2.82 (2H, m), 3.04 (1H, m), 3.33 (1H, m), 3.48 (6H, s), 3.56-3.67 (2H, m), 6.55 (1H, s), 7.03 (1H, d, J=8.4 Hz), 7.30-7.56 (6H, m), 7.78-7.85 (6H, m), 8.04 (2H, d, J=8.4 Hz), 10.17 (1H, s).
- 30

FABMS(pos) 419.1 [M+H]⁺

Example 173

N-[6-[(Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-sulfonamide
hydrochloride



6-[(N, N-Dimethylamino)methyl]-7,8-dihydro-2-naphthalenamine (200 mg, 0.72 mmol) obtained in Example 41-2) was dissolved in acetonitrile (30 ml). Triethylamine (0.401 ml, 2.88 mmol) and [1,1'-biphenyl]-4-sulfonylchloride (200 mg, 0.79 mmol) were added to the solution under ice-cooling, which was stirred for 3 hours. The reaction mixture was concentrated. Ethyl acetate and water were added to the residue, and extraction was conducted. The ethyl acetate layer was concentrated, and the residue was purified by alumina column chromatography (development solvent; ethyl acetate:n-hexane = 33:67). 4N Hydrogen chloride-ethyl acetate solution was added to the resulting oily substance, which was concentrated. The residue was recrystallized from methanol - ethyl acetate, to give the titled compound (194 mg).

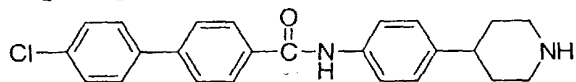
¹H-NMR (DMSO-d₆) δ: 1.32 (1H, m), 1.96 (1H, m), 2.11 (1H, m), 2.35 (1H, d, J=15.9, 10.0 Hz), 2.74 (2H, m), 2.78 (7H, m), 3.02 (2H, m), 6.89 (2H, d, J=10.6 Hz), 6.91 (1H, m), 7.40-7.51 (3H, m), 7.70 (2H, d, J=6.7 Hz), 7.85 (4H, m), 9.92 (1H, brs), 10.23 (1H, s).

Melting point: 168-170 °C (crystallization solvent: methanol - ethyl acetate).

FABMS(pos) 421.1 [M+H]⁺

Example 174

4'-Chloro-N-[4-(4-piperidinyl)phenyl][1,1'-biphenyl]-4-carboxamide

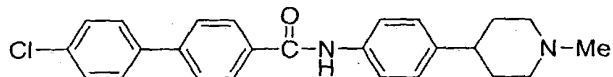


5 The titled compound was obtained as a colorless powder by carrying out the same operation as in Example 127-2), using 4'-chloro-N-[4-(4-piperidinyl)phenyl][1,1'-biphenyl]-4-carboxamide obtained in Reference Example 89. ¹H-NMR (CDCl₃+ DMSO-d₆) δ: 1.40-1.90 (4H, m), 2.60-2.90 (3H, m), 3.18-3.28 (2H, m), 7.19 (2H, d, J=8.1 Hz), 7.49 (2H, d, J=7.0 Hz), 7.67-7.75 (6H, m), 8.07-8.10 (3H, m), 10.16 (1H, s).
Melting point: 276-281 °C (decomposition) (crystallization solvent: ethyl acetate)

15

Example 175

4'-Chloro-N-[4-(1-methyl4-piperidinyl)phenyl][1,1'-biphenyl]-4-carboxamide



20 A mixture of 4'-chloro-N-[4-(4-piperidinyl)phenyl][1,1'-biphenyl]-4-carboxamide (0.17 g) obtained in Example 174, 37% aqueous formaldehyde solution (0.05 ml) and formic acid (0.5 ml) was heated at 100°C for 4 hours. The reaction mixture was cooled to room
25 temperature. Water was added to the mixture, which was made alkaline with 8N aqueous sodium hydroxide solution, and extracted with ethyl acetate - tetrahydrofuran (1:1) mixed solution. The extract was washed with saturated aqueous sodium chloride solution, dried over anhydrous magnesium
30 sulfate, and then the solvent was distilled out under reduced pressure. The resulting solid was washed with ethyl acetate, dried under reduced pressure, to give the titled compound (90 mg).

¹H-NMR (CDCl₃+ DMSO-d₆) δ: 1.55-1.80 (2H, m), 1.90-2.10

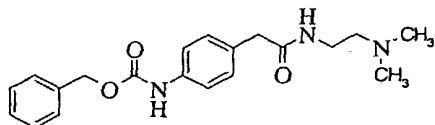
(2H, m), 2.22 (3H, s), 2.30-2.45 (1H, m), 2.80-3.20 (4H, m), 7.11 (2H, d, J=8.1 Hz), 7.36 (2H, d, J=8.1 Hz), 7.50-7.63 (6H, m), 7.97 (2H, d, J=8.4 Hz), 9.79 (1H, s).

Melting point: 273-277 °C (decomposition) (Washing

5 solvent: ethyl acetate)

Example 176

Benzyl 4-[2-[[2-(dimethylamino)ethyl]amino]-2-oxoethyl]phenylcarbamate



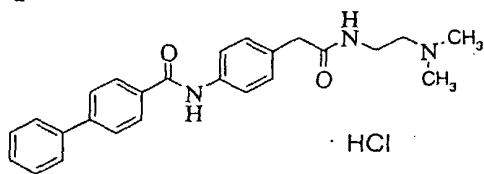
10

N,N-Dimethylethylenediamine (0.64 ml), WSC (1.31 g), HOBT (1.05 g), and triethylamine (2.4 ml) were added to a tetrahydrofuran (50 ml) solution of 2-[4-
 15 [[(benzyloxy)carbonyl]amino]phenyl]acetic acid (1.5 g) obtained in Reference Example 90. After stirring for 20 hours, the reaction mixture was poured into water, and extraction was conducted using ethyl acetate. The organic layer was washed with water, saturated aqueous sodium bicarbonate solution, and saturated aqueous sodium
 20 chloride solution, dried and then concentrated. The residue was recrystallized from ethyl acetate - hexane, to give the titled compound (1.72 g).

Melting point: 126-127 °C.

25 Example 177

N-[4-[2-[[2-(Dimethylamino)ethyl]amino]-2-oxoethyl]phenyl][1,1'-biphenyl]-4-carboxamide hydrochloride



30

Oxalyl chloride (0.56 ml) was added dropwise to a

tetrahydrofuran (45 ml) solution of 4-biphenylcarboxylic acid (1.01 g) under ice-cooling. 9 drops of DMF was added to the mixture, and the temperature of the mixture was raised to room temperature, which was stirred for 40 minutes. The reaction mixture was concentrated and dried.

A tetrahydrofuran (50 ml) solution of the residue was added dropwise to a tetrahydrofuran (45 ml) solution of 2-(4-aminophenyl)-N-[2-(dimethylamino)ethyl]acetamide (939 mg) obtained in Reference Example 91 under ice-cooling. Then the temperature of the reaction mixture was raised to room temperature, which was stirred for 2 hours. Saturated aqueous sodium bicarbonate solution was added to the reaction mixture, and extraction was conducted using ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried over sodium sulfate, and then concentrated. The residue was dissolved in tetrahydrofuran. 4N Hydrochloric acid-ethyl acetate was added to the solution, which was concentrated.

The residue was recrystallized from methanol - diisopropyl ether, to give the titled compound (750 mg). Melting point: 216-217 °C.

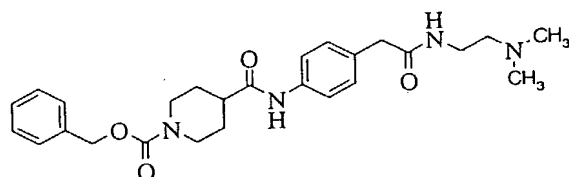
The above N-[4-[2-[[2-(dimethylamino)ethyl]amino]-2-oxoethyl]phenyl][1,1'-biphenyl]-4-carboxamide hydrochloride (100 mg) was dissolved in saturated aqueous sodium bicarbonate solution, and extraction was conducted using tetrahydrofuran-ethyl acetate (1:1). The organic layer was washed with saturated aqueous sodium chloride solution, dried over sodium sulfate, and then concentrated.

The residue was recrystallized from methanol - diisopropyl ether, to give a free base form (56 mg) of the titled compound.

Melting point: 228-229 °C.

Example 178

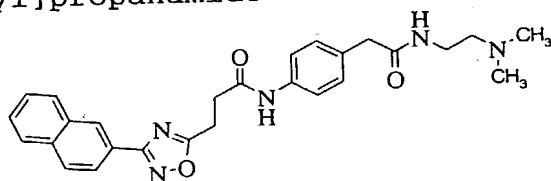
Benzyl 4-[[4-[2-[[2-(dimethylamino)ethyl]amino]-2-oxoethyl]anilino]carbonyl]-1-piperidinecarboxylate



2-(4-Aminophenyl)-N-[2-(dimethylamino)ethyl]acetamide (221 mg), WSC (249 mg), 1-hydroxybenzotriazole (199 mg), triethylamine (0.4 ml), and dimethylaminopyridine (244 mg) were added to a tetrahydrofuran (10 ml) solution of 1-[(benzyloxy)carbonyl]-4-piperidinecarboxylic acid (290 mg), which was stirred for 20 hours. The reaction mixture was poured into water, and extraction was conducted using ethyl acetate. The organic layer was washed with water, saturated aqueous sodium bicarbonate solution, and saturated aqueous sodium chloride solution, dried over sodium sulfate, and then concentrated. The residue was recrystallized from methanol - diisopropyl ether, to give the titled compound (230 mg).
Melting point: 169-170 °C.

Example 179

N-[4-[2-[[2-(Dimethylamino)ethyl]amino]-2-oxoethyl]phenyl]-3-[3-(2-naphthyl)-1,2,4-oxadiazol-5-yl]propanamide

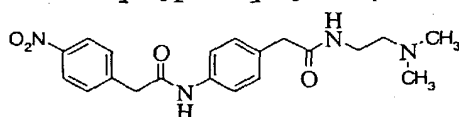


2-(4-Aminophenyl)-N-[2-(dimethylamino)ethyl]acetamide (221 mg), WSC (249 mg), 1-hydroxybenzotriazole (199 mg), triethylamine (0.4 ml), and dimethylaminopyridine (244 mg) were added to a DMF (5 ml) solution of 3-[3-(2-naphthyl)-1,2,4-oxadiazol-5-yl]propionic acid (268 mg), which was stirred for 5 hours. The reaction mixture was poured into water, and extraction was conducted using ethyl acetate. The organic layer was

washed with water, saturated aqueous sodium bicarbonate solution, and saturated aqueous sodium chloride solution, dried over sodium sulfate, and then concentrated. The residue was recrystallized from methanol, to give the
 5 titled compound (166 mg).
 Melting point: 173-174 °C.

Example 180

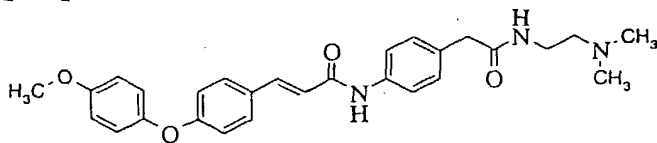
N-[4-[2-[[2-(Dimethylamino)ethyl]amino]-2-
 10 oxoethyl]phenyl]-2-(4-nitrophenyl)acetamide



2-(4-Aminophenyl)-N-[2-(
 (dimethylamino)ethyl]acetamide (221 mg), WSC (free form :
 0.23 ml), 1-hydroxybenzotriazole (199 mg), and
 15 dimethylaminopyridine (244 mg) were added to a DMF (5 ml)
 solution of 4-nitrophenylacetic acid (181 mg), which was
 stirred for 4 hours. The reaction mixture was poured into
 water, and extraction was conducted using ethyl acetate.
 The organic layer was washed with water, saturated aqueous
 20 sodium bicarbonate solution, and saturated aqueous sodium
 chloride solution, dried over sodium sulfate, and then
 concentrated. The residue was recrystallized from
 methanol, to give the titled compound (80 mg).
 Melting point: 160-162 °C.

Example 181

(E)-N-[4-[2-[[2-(Dimethylamino)ethyl]amino]-2-
 oxoethyl]phenyl]-3-[4-(4-methoxyphenoxy)phenyl]-2-
 30 propanamide

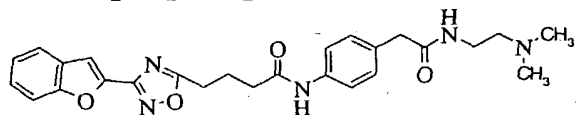


2-(4-Aminophenyl)-N-[2-(
 (dimethylamino)ethyl]acetamide (221 mg), WSC (free form :

0.23 ml), 1-hydroxybenzotriazole (199 mg), triethylamine (0.14 ml) and dimethylaminopyridine (122 mg) were added to a DMF (5 ml) solution of (E)-3-[4-(4-methoxyphenoxy)phenyl]-2-propenoic acid (270 mg), which
 5 was stirred for 24 hours. The reaction mixture was poured into water, and extraction was conducted using ethyl acetate - tetrahydrofuran (1:1). The organic layer was washed with water, saturated aqueous sodium bicarbonate solution, and saturated aqueous sodium chloride solution,
 10 dried over sodium sulfate, and then concentrated. The resulting crude crystals were washed with diisopropyl ether, to give the titled compound (227 mg).
 Melting point: 175-177 °C (decomposition).

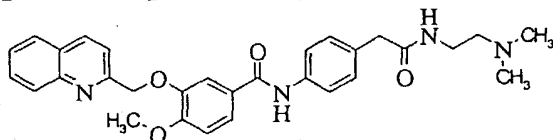
15 Compounds described in the following Example 182 to 198 were produced in the same manner as in Example 181.
 Example 182

4-[3-(1-Benzofuran-2-yl)-1,2,4-oxadiazol-5-yl]-N-[4-[2-
 20 [[2-(dimethylamino)ethyl]amino]-2-oxoethyl]phenyl]butanamide



Melting point: 161-163 °C.
 Washing solvent: diisopropyl ether.

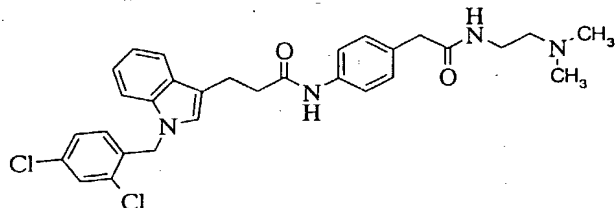
25 Example 183
 N-[4-[2-[[2-(Dimethylamino)ethyl]amino]-2-oxoethyl]phenyl]-3-methoxy-4-(2-quinolinylmethoxy)benzamide



30 Melting point: 209-210 °C (decomposition).
 Washing solvent: diisopropyl ether.

Example 184

3-[1-(2,4-Dichlorobenzyl)-1H-indol-3-yl]-N-[4-[2-[[2-(dimethylamino)ethyl]amino]-2-oxoethyl]phenyl]propanamide



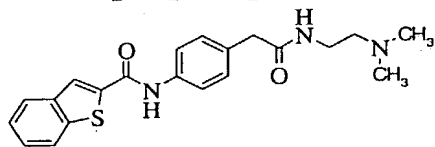
5

Melting point: : 123-125 °C (decomposition).

Washing solvent: diisopropyl ether.

Example 185

10 N-[4-[2-[[2-(Dimethylamino)ethyl]amino]-2-oxoethyl]phenyl]-1-benzothiophen-2-carboxamide



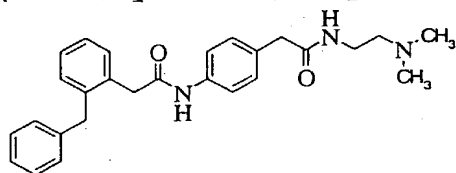
Melting point: 186-187 °C (decomposition).

Washing solvent: diisopropyl ether.

15

Example 186

2-(2-Benzylphenyl)-N-[4-[2-[[2-(dimethylamino)ethyl]amino]-2-oxoethyl]phenyl]acetamide

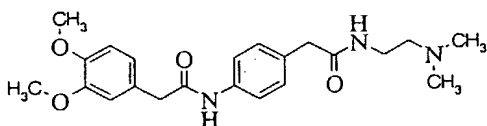


20 Melting point: 115-117 °C.

Washing solvent: diisopropyl ether.

Example 187

25 2-(3,4-dimethoxyphenyl)-N-[4-[2-[[2-(dimethylamino)ethyl]amino]-2-oxoethyl]phenyl]acetamide

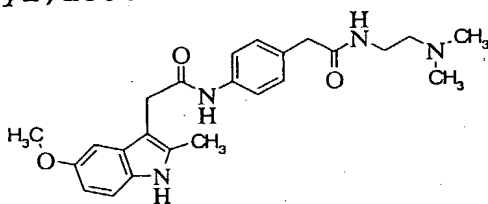


Melting point: 123-124 °C.

Recrystallization solvent: methanol - diisopropyl ether.

5 Example 188

N-[4-[2-[[2-(Dimethylamino)ethyl]amino]-2-oxoethyl]phenyl]-2-(5-methoxy-2-methyl-1H-indol-3-yl)acetamide

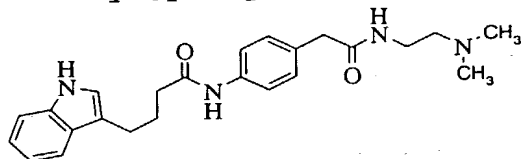


10 Melting point: 125-126 °C.

Recrystallization solvent: methanol - diisopropyl ether.

Example 189

15 N-[4-[2-[[2-(Dimethylamino)ethyl]amino]-2-oxoethyl]phenyl]-4-(1H-indol-3-yl)butanamide

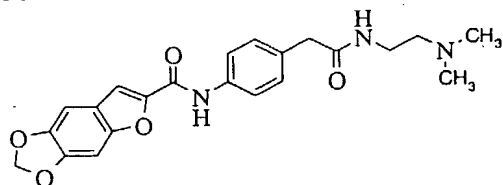


Melting point: 132-133 °C.

Washing solvent: diisopropyl ether.

20 Example 190

N-[4-[2-[[2-(Dimethylamino)ethyl]amino]-2-oxoethyl]phenyl]furo[2,3-f][1,3]benzodioxol-6-carboxamide

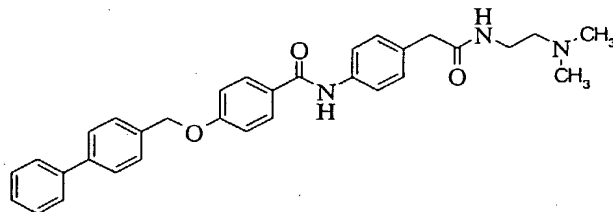


Melting point: : 173-175 °C (decomposition).

Washing solvent: diisopropyl ether.

Example 191

- 5 4-([1,1'-Biphenyl]-4-ylmethoxy)-N-[4-[2-[[2-(dimethylamino)ethyl]amino]-2-oxoethyl]phenyl]benzamide



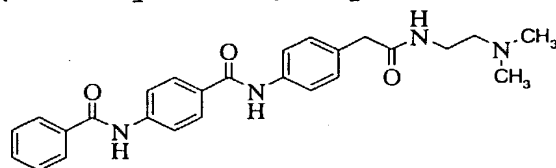
Melting point: 204-208 °C.

Washing solvent: diisopropyl ether.

10

Example 192

- 4-(Benzoylamino)-N-[4-[2-[[2-(dimethylamino)ethyl]amino]-2-oxoethyl]phenyl]benzamide

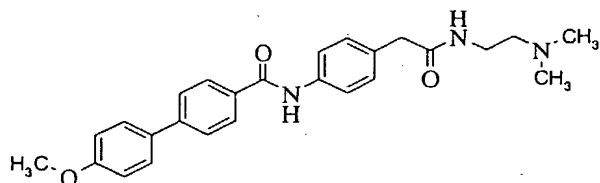


- 15 Melting point: 220-221 °C.

Washing solvent: diisopropyl ether.

Example 193

- 20 N-[4-[2-[[2-(Dimethylamino)ethyl]amino]-2-oxoethyl]phenyl]-4'-methoxy[1,1'-biphenyl]-4-carboxamide



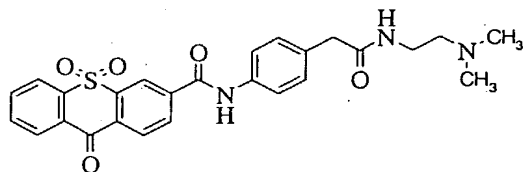
Melting point: 196-198 °C (decomposition).

Washing solvent: diisopropyl ether.

25

Example 194

N-[4-[2-[[2-(Dimethylamino)ethyl]amino]-2-oxoethyl]phenyl]-9,10,10-trioxo-9,10-dihydro-10 λ^6 -thioxanten-3-carboxamide



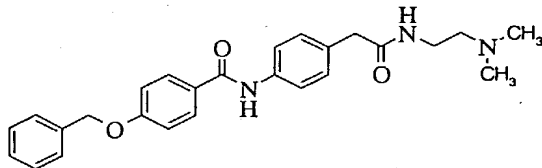
5

Melting point: 162-163 °C (decomposition).

Washing solvent: diisopropyl ether.

Example 195

4-(Benzyloxy)-N-[4-[2-[[2-(dimethylamino)ethyl]amino]-2-oxoethyl]phenyl]benzamide

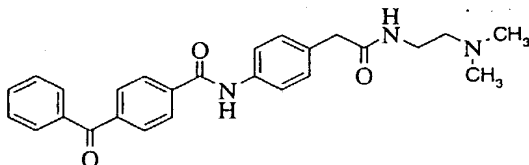


Melting point: 190-192 °C (decomposition).

15 Washing solvent: diisopropyl ether.

Example 196

4-Benzoyl-N-[4-[2-[[2-(dimethylamino)ethyl]amino]-2-oxoethyl]phenyl]benzamide



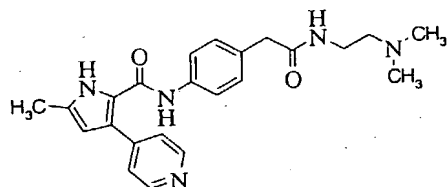
20

Melting point: 173-175 °C (decomposition).

Washing solvent: diisopropyl ether.

Example 197

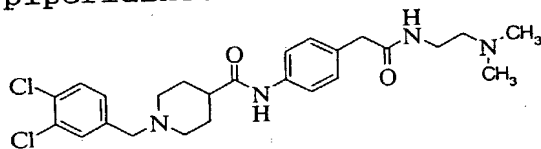
25 N-[4-[2-[[2-(Dimethylamino)ethyl]amino]-2-oxoethyl]phenyl]-5-methyl-3-(4-pyridinyl)-1H-pyrrole-2-carboxamide



Melting point: : 215-218 °C (decomposition).
 Washing solvent: diisopropyl ether.

5 Example 198

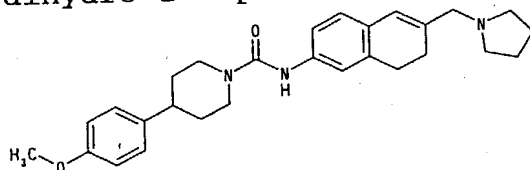
1-(3,4-Dichlorobenzyl)-N-[4-[2-[[2-(dimethylamino)ethyl]amino]-2-oxoethyl]phenyl]-4-piperidinecarboxamide



10 Melting point: : 182-183 °C (decomposition).
 Washing solvent: diisopropyl ether.

Example 199

15 4-(4-Methoxyphenyl)-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide



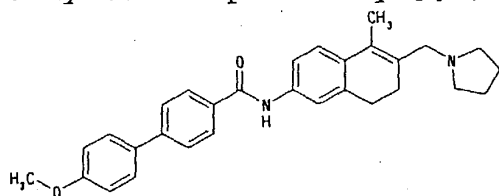
The titled compound was obtained by carrying out the same operation as in Example 99, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

20 ¹H-NMR (CDCl₃) δ: 1.61-1.91 (8H, m), 2.31 (2H, t, J=8.1 Hz), 2.54 (4H, m), 2.73-2.81 (3H, m), 2.98 (2H, t, J=7.8 Hz), 3.16 (2H, s), 3.79 (3H, s), 4.20 (2H, d, J=13.1 Hz), 6.31 (1H, s), 6.36 (1H, s), 6.86 (2H, d, J=8.6 Hz), 7.06-7.20 (5H, m).

25 Melting point: 175-176 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 200

4'-Methoxy-N-[6-(1-pyrrolidinylmethyl)-5-methyl-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



5

The titled compound was obtained by carrying out the same operation as in Example 1, using 5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 69.

10 $^1\text{H-NMR}$ (CDCl_3) δ : 1.78 (4H, m), 2.10 (3H, s), 2.37 (2H, t, $J=8.1$ Hz), 2.53 (4H, m), 2.76 (2H, t, $J=8.1$ Hz), 3.28 (2H, s), 3.87 (3H, s), 7.01 (2H, d, $J=8.6$ Hz), 7.27 (1H, d, $J=7.8$ Hz), 7.46 (1H, d, $J=7.8$ Hz), 7.48 (1H, s), 7.57 (2H, d, $J=8.6$ Hz), 7.66 (2H, d, $J=8.4$ Hz), 7.81 (1H, s), 7.92 (2H, d, $J=8.4$ Hz).

15

Elemental analysis for $\text{C}_{30}\text{H}_{32}\text{N}_2\text{O}_2$

Calcd.: C, 79.61; H, 7.13; N, 6.19

Found: C, 79.35; H, 7.28; N, 6.24

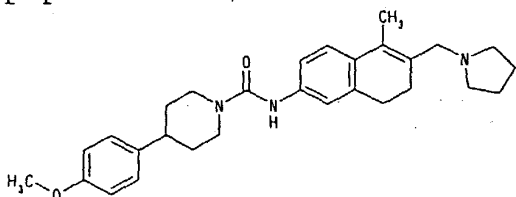
Melting point: 179-180 $^\circ\text{C}$ (crystallization solvent: ethyl acetate - diisopropyl ether)

20

Example 201

4-(4-Methoxyphenyl)-N-[6-(1-pyrrolidinylmethyl)-5-methyl-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide

25



The titled compound was obtained by carrying out the same operation as in Example 99, using 5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine

obtained in Reference Example 69.

$^1\text{H-NMR}$ (CDCl_3) δ : 1.67 (2H, dd, $J=13.4$, 4.0 Hz), 1.78 (4H, m), 1.89 (2H, d, $J=11.4$ Hz), 2.07 (3H, s), 2.34 (2H, t, $J=7.5$ Hz), 2.52 (4H, m), 2.68-2.73 (3H, m), 2.98 (2H, t, $J=7.5$ Hz),
 5 3.26 (2H, s), 3.80 (3H, s), 4.20 (2H, d, $J=13.4$ Hz), 6.36 (1H, s), 6.86 (2H, d, $J=8.4$ Hz), 7.12-7.20 (5H, m).

Elemental analysis for $\text{C}_{28}\text{H}_{37}\text{N}_3\text{O}_2$

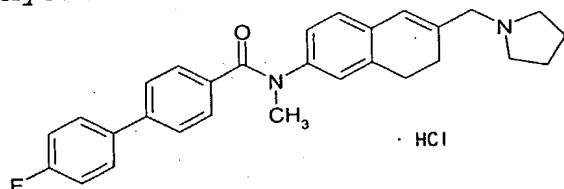
Calcd.: C, 75.13; H, 8.33; N, 9.39

Found: C, 74.96; H, 8.14; N, 9.10

10 Melting point: 163-164 $^\circ\text{C}$ (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 202

15 4'-Fluoro-N-methyl-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide hydrochloride



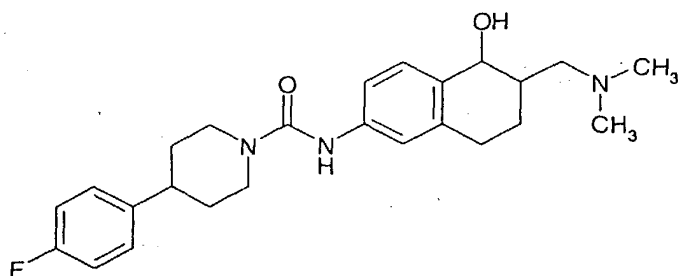
The titled compound was obtained by carrying out the same operation as in Example 1, using N-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine hydrochloride obtained in Reference Example 95.

20 $^1\text{H-NMR}$ ($\text{DMSO}-d_6$) δ : 1.92-1.98 (4H, m), 2.39 (2H, t, $J=8.1$ Hz), 2.73 (2H, t, $J=8.1$ Hz), 3.00 (2H, m), 3.35 (3H, m), 3.44 (2H, m), 3.83 (2H, d, $J=5.6$ Hz), 6.62 (1H, s), 6.92-7.01
 25 (2H, m), 7.11 (1H, s), 7.26 (2H, dd, $J=8.9$, 5.6 Hz), 7.38 (2H, d, $J=8.1$ Hz), 7.55 (2H, d, $J=8.1$ Hz), 7.69 (2H, dd, $J=8.9$, 5.6 Hz), 10.60 (1H, brs).

FABMS(pos) 441.2 $[\text{M}+\text{H}]^+$

30 Example 203

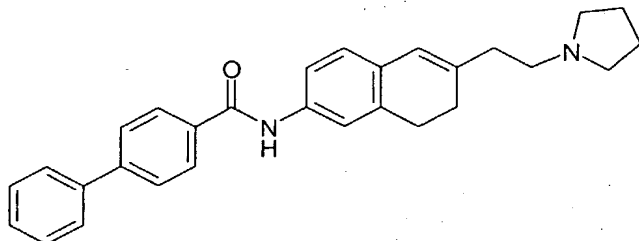
N-[6-[(Dimethylamino)methyl]-5-hydroxy-5,6,7,8-tetrahydro-2-naphthalenyl]-4-(4-fluorophenyl)-1-piperidinecarboxamide



N,N-Dimethylmethylethylamine hydrochloride (638 mg, 6.82 mmol) was added to a mixed solution of 4-(4-fluorophenyl)-N-(5-oxo-5,6,7,8-tetrahydro-2-naphthalenyl)-1-piperidinecarboxamide (1.00 g, 2.73 mmol) obtained in Reference Example 97 in tetrahydrofuran (10 ml) and acetonitrile (10 ml), which was stirred at room temperature for 1 day. The solvent was distilled out under reduced pressure. Ethyl acetate was added to the residue, which was washed with aqueous potassium carbonate solution and saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. The resulting oily substance was dissolved in methanol (15 ml). Sodium borohydride (103 mg, 2.73 mmol) was added to the solution under ice-cooling, which was stirred for 1 hour. Then, the solvent was distilled out under reduced pressure. 1N Hydrochloric acid was added to the residue, which was washed with ethyl acetate. 4N Sodium hydroxide was added to the water layer to make it alkaline. The reaction mixture was extracted with ethyl acetate, which was washed with saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. The resulting residue was purified by aluminum B column chromatography (development solvent; ethyl acetate), powdered with hexane, to give the titled compound (231 mg). Melting point: 160-163 °C (crystallization solvent: ethyl acetate - n-hexane) FAB(pos) 426.3 [M+H]+

Example 204

N-[6-[2-(1-Pyrrolidinyl)ethyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



- 5 Concentrated hydrochloric acid (2 ml) was added to N-[6-[2-(1-pyrrolidinyl)ethyl]-7,8-dihydro-2-naphthalenyl]acetamide (98.0 mg, 0.345 mmol) obtained in Reference Example 103, which was stirred at 100 °C for 16 hours. The solvent was distilled out under reduced
- 10 pressure. Ethyl acetate was added to the residue, which was washed with aqueous potassium carbonate solution and saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. WSC (62.5 mg, 0.326
- 15 mmol) was added to a dimethylformamide solution (1.5ml) of the resulting oily substance (79.0 mg, 0.326 mmol), [1,1'-biphenyl]-4-carboxylic acid (64.6 mg, 0.326 mmol) and DMAP (39.8 mg, 0.326 mmol) under ice-cooling, which was stirred at room temperature for 1 day. Ethyl acetate was
- 20 added to the reaction mixture, washed with aqueous potassium carbonate solution and saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, then the solvent was distilled out under reduced pressure. The resulting residue was purified by aluminum column
- 25 chromatography (development solvent; ethyl acetate), powdered with ethyl acetate and isopropyl ether (1:5), to give the titled compound (36.8 mg).
- ¹H NMR (DMSO-d₆) δ: 1.67 (4H, m), 2.23 (2H, m), 2.34 (2H, m), 2.46 (4H, m), 2.57 (2H, m), 2.75 (2H, m), 6.24 (1H, s),
- 30 6.98 (1H, d, J = 8.1 Hz), 7.40-7.59 (5H, m), 7.76 (2H, d, J = 7.5 Hz), 7.82 (2H, d, J=8.4 Hz), 8.05 (2H, d, J = 8.4

Hz), 10.19 (1H, s).

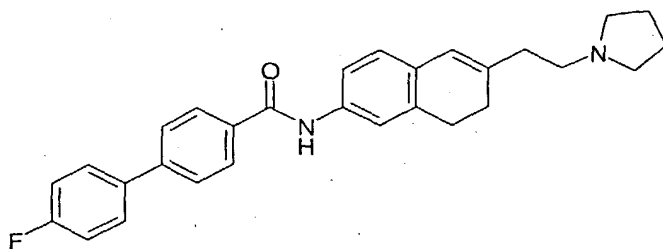
Meltingpoint: 184-186 °C (crystallization solvent: ethyl acetate - isopropyl ether)

FAB(pos) 423.2 [M+H]⁺

5

Example 205

4'-Fluoro-N-[6-[2-(1-pyrrolidinyl)ethyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



- 10 Concentrated hydrochloric acid (2 ml) was added to N-[6-[2-(1-pyrrolidinyl)ethyl]-7,8-dihydro-2-naphthalenyl]acetamide (98.0 mg, 0.345 mmol) obtained in Reference Example 103, which was stirred at 100°C for 16 hours. The solvent was distilled out under reduced
- 15 pressure. Ethyl acetate was added to the residue, which was washed with aqueous potassium carbonate solution and saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. WSC (62.5 mg, 0.326
- 20 mmol) was added to a dimethylformamide solution (1.5 ml) of the resulting oily substance (79.0 mg, 0.326 mmol), 4'-fluoro-[1,1'-biphenyl]-4-carboxylic acid (64.6 mg, 0.326 mmol) and DMAP (39.8 mg, 0.326 mmol) under ice-cooling, which was stirred at room temperature for 1 day.
- 25 Ethyl acetate was added to the reaction mixture, which was washed with aqueous potassium carbonate solution and saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and the solvent was distilled out under reduced pressure. The resulting residue was
- 30 purified by aluminum column chromatography (development solvent; ethyl acetate), powdered with ethyl acetate -

isopropyl ether (1:5) , to give the titled compound (75.1 mg).

¹H NMR (DMSO-d₆) δ: 1.68 (4H, m), 2.23 (2H, m), 2.35 (2H, m), 2.50 (4H, m), 2.59 (2H, m), 2.75 (2H, m), 6.24 (1H, s),
5 6.98 (1H, d, J = 8.1 Hz), 7.34 (2H, m), 7.56 (2H, m), 7.81 (4H, m), 8.04 (2H, d, J = 8.4 Hz), 10.19 (1H, s).

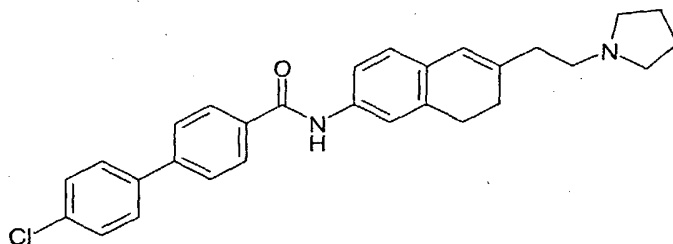
Melting point: 187-189°C (crystallization solvent: ethyl acetate - isopropyl ether)

FAB (pos) 441.3 [M+H]⁺

10

Example 206

4'-Chloro-N-[6-[2-(1-pyrrolidinyl)ethyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



15 Concentrated hydrochloric acid (2 ml) was added to N-[6-[2-(1-pyrrolidinyl)ethyl]-7,8-dihydro-2-naphthalenyl]acetamide (98.0 mg, 0.345 mmol) obtained in Reference Example 103, which was stirred at 100°C for 16 hours. The solvent was distilled out under reduced
20 pressure. Ethyl acetate was added to the residue, which was washed with aqueous potassium carbonate solution and saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. WSC (62.5 mg, 0.326
25 mmol) was added to a dimethylformamide solution (1.5 ml) of the resulting oily substance (79.0 mg, 0.326 mmol), 4'-chloro-[1,1'-biphenyl]-4-carboxylic acid (64.6 mg, 0.326 mmol) and DMAP (39.8 mg, 0.326 mmol) under ice-cooling, which was stirred at room temperature for 1 day.
30 Ethyl acetate was added to the reaction mixture, which was washed with aqueous potassium carbonate solution and

saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. The resulting residue was purified by aluminum column chromatography (development solvent; ethyl acetate), powdered with ethyl acetate - isopropyl ether (1:5), to give the titled compound (78.4 mg).

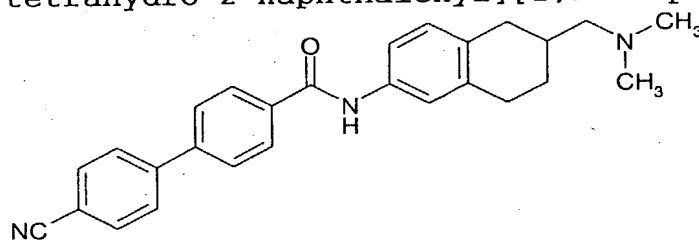
¹H NMR (DMSO-d₆) δ: 1.67 (4H, m), 2.23 (2H, m), 2.34 (2H, m), 2.45 (4H, m), 2.57 (2H, m), 2.75 (2H, m), 6.24 (1H, s), 6.98 (1H, d, J = 8.1 Hz), 7.55 (4H, m), 7.80 (2H, d, J=8.4 Hz), 7.84 (2H, d, J=8.4 Hz), 8.05 (2H, d, J = 8.7 Hz), 10.20 (1H, s).

Melting point: 207-209°C (crystallization solvent: ethyl acetate - isopropyl ether)

FAB (pos) 457.2 [M+H]⁺

Example 207

4'-Cyano-N-[6-[(dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the same operation as in Example 1, using N-[(6-amino-1,2,3,4-tetrahydro-2-naphthalenyl)methyl]-N,N-dimethylamine and 4'-cyano-[1,1'-biphenyl]-4-carboxylic acid.

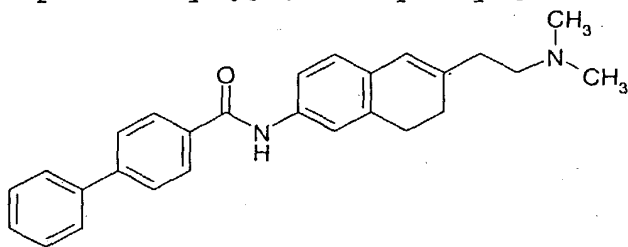
¹H NMR (CDCl₃) δ: 1.42 (1H, m), 1.95 (2H, m), 2.26 (6H, s), 2.24-2.46 (3H, m), 2.84-2.95 (3H, m), 7.10 (1H, d, J=8.4 Hz), 7.30 (1H, m), 7.46 (1H, s), 7.74 (7H, m), 7.98 (2H, d, J=8.4 Hz).

Melting point: 183-185°C (crystallization solvent: ethyl acetate - isopropyl ether)

FAB (pos) 410.2 [M+H]⁺

Example 208

N-[6-[2-(Dimethylamino)ethyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



5 Concentrated hydrochloric acid (1.5 ml) was added to N-[6-[2-(dimethylamino)ethyl]-7,8-dihydro-2-naphthalenyl]acetamide (57.5 mg, 0.223 mmol) obtained in Reference Example 104, which was stirred at 100°C for 1
10 hour. The solvent was distilled out under reduced pressure. Ethyl acetate was added to the residue, which was washed with aqueous potassium carbonate solution and saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was
15 distilled out under reduced pressure. WSC (29.2 mg, 0.139 mmol) was added to a dimethylformamide solution (0.7 ml) of the resulting oily substance (30 mg, 0.139 mmol), [1,1'-biphenyl]-4-carboxylic acid (30.2 mg, 0.139 mmol) and DMAP (16.9 mg, 0.139 mmol) under ice-cooling, which was
20 stirred at room temperature for 16 hours. Ethyl acetate was added to the reaction mixture, which was washed with aqueous potassium carbonate solution and saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under
25 reduced pressure. The resulting residue was purified by aluminum column chromatography (development solvent; ethyl acetate), powdered with ethyl acetate - isopropyl ether (1:5), to give the titled compound (12.4 mg).
¹H NMR (CDCl₃) δ: 2.29 (8H, m), 2.41 (2H, m), 2.46 (2H, m),
30 2.84 (2H, t, J = 8.1 Hz), 6.24 (1H, s), 6.98 (1H, d, J = 8.4 Hz), 7.34 (1H, m), 7.41 (1H, d, J = 6.9 Hz), 7.46 (3H,

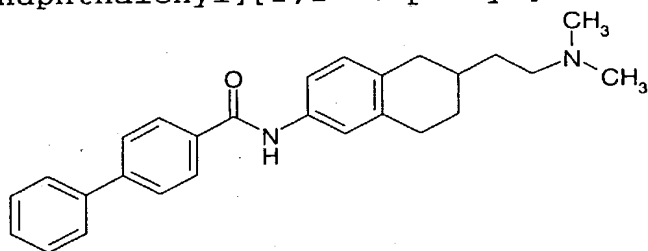
m), 7.63 (2H, d, J = 7.2 Hz), 7.71 (2H, d, J = 8.4 Hz), 7.77 (1H, br), 7.94 (2H, d, J = 8.4 Hz).

Melting point: 148-150°C (crystallization solvent: ethyl acetate - isopropyl ether)

5 FAB (pos) 397.2 [M+H]⁺

Example 209

N-[6-[2-(Dimethylamino)ethyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



10

A methanol solution (5 ml) of N-[6-[2-(dimethylamino)ethyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide (20 mg, 0.050 mmol) obtained in Example 208 and palladium carbon (10 mg) was stirred under hydrogen atmosphere for 4 hours. After a catalyst was filtered off, the filtrate was concentrated under reduced pressure. The resulting residue was purified by aluminum B column chromatography (development solvent; ethyl acetate), powdered with ethyl acetate - hexane (1:3), to give the titled compound (4.0 mg).

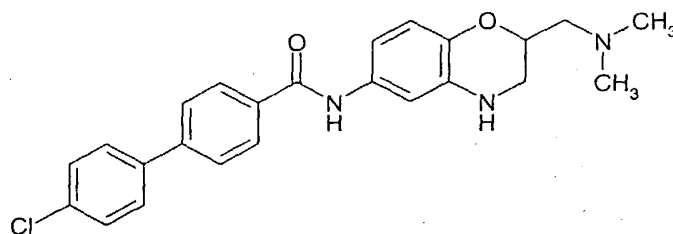
¹H NMR (CDCl₃) δ: 1.60 (4H, m), 1.92 (1H, m), 2.26 (6H, s), 2.42 (3H, m), 2.84 (3H, m), 7.06 (1H, d, J=8.1Hz), 7.32 (1H, m), 7.46 (4H, m), 7.63 (2H, d, J=6.9Hz), 7.72 (3H, m), 7.94 (2H, d, J=8.1Hz).

25 Melting point: 112-114°C (crystallization solvent: ethyl acetate - isopropyl ether)

FAB(pos) 399.2 [M+H]⁺

Example 210

30 4'-Chloro-N-[2-[(dimethylamino)methyl]-3,4-dihydro-2H-1,4-benzoxazin-6-yl][1,1'-biphenyl]-4-carboxamide



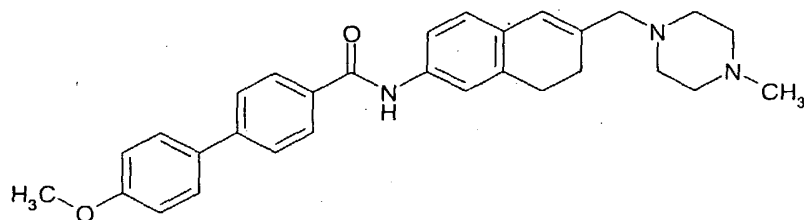
The titled compound was obtained as white powders by the same method as in Example 1, using 6-amino-2-(dimethylamino)methyl-1,4-benzoxazin obtained in Reference Example 105.

¹H-NMR (CDCl₃) δ: 2.33 (6H, s), 2.44-2.65 (2H, m), 3.15-3.21 (1H, m), 3.41-3.46 (1H, m), 3.87 (1H, brs), 4.24-4.26 (1H, m), 6.61 (1H, dd, J=2.5, 8.6 Hz), 6.81 (1H, d, J=8.6 Hz), 7.28 (1H, d, J=2.5 Hz), 7.43 (2H, d, J=6.5 Hz), 7.54 (2H, d, J=6.5 Hz), 7.64 (2H, d, J=8.4 Hz), 7.71 (1H, s), 7.90 (2H, d, J=8.4 Hz).

Melting point: 227-230 °C (crystallization solvent: diisopropyl ether)

Example 211

4'-Methoxy-N-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



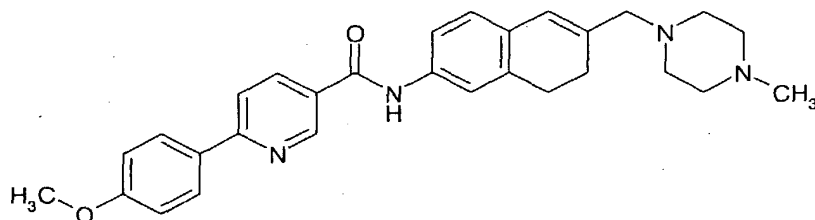
The titled compound was obtained as colorless powders by carrying out the same operation as in Example 1, using 6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenamine obtained in Reference Example 106.

¹H NMR (CDCl₃) δ: 2.31 (3H, s), 2.33 (2H, t, J=8.1 Hz), 2.49 (8H, bs), 2.84 (2H, t, J=8.1 Hz), 3.07 (2H, s), 3.87 (3H, s), 6.36 (1H, s), 7.00-7.03 (3H, m), 7.36 (1H, d, J=8.1 Hz), 7.51 (1H, s), 7.58 (2H, d, J=8.4 Hz), 7.67 (2H, d, J=8.4 Hz), 7.78 (1H, s), 7.91 (2H, d, J=8.4 Hz).

Melting point: 208-210 °C (crystallization solvent: ethyl acetate)

Example 212

5 6-(4-Methoxyphenyl)-N-[6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl]nicotinamide



10 The titled compound was obtained as colorless powders by carrying out the same operation as in Example 1, using 6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenamine obtained in Reference Example 106.

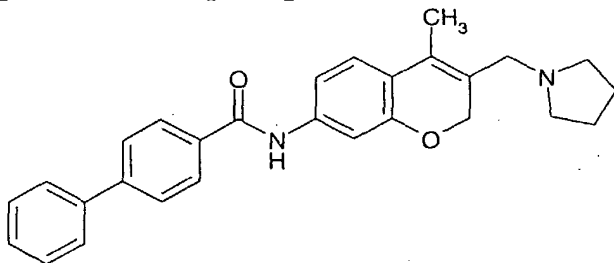
¹H NMR (CDCl₃) δ: 2.30 (3H, s), 2.33 (2H, t, J=8.1 Hz), 2.47 (8H, bs), 2.84 (2H, t, J=8.1 Hz), 3.07 (2H, s), 3.89 (3H, s), 6.36 (1H, s), 7.01-7.04 (3H, m), 7.37 (1H, d, J=8.1 Hz), 7.49 (1H, s), 7.78-7.81 (2H, m), 8.03 (2H, d, J=8.4 Hz), 8.21 (1H, dd, J=2.1 Hz, 8.7 Hz), 9.09 (1H, s).

Melting point: 235-237 °C (crystallization solvent: ethyl acetate)

20

Example 213

N-[4-Methyl-3-(1-pyrrolidinylmethyl)-2H-chromen-7-yl][1,1'-biphenyl]-4-carboxamide



25

The titled compound was obtained as colorless powders by carrying out the same operation as in Example 1, using

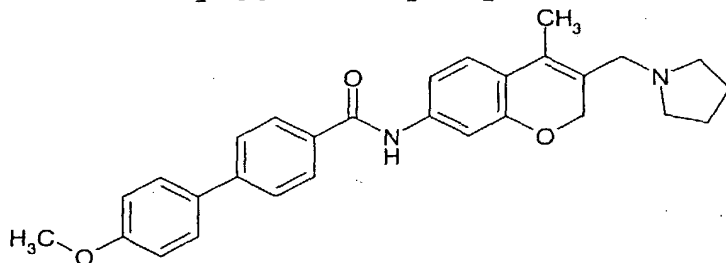
4-methyl-3-(1-pyrrolidinymethyl)-2H-chromen-7-amine
obtained in Reference Example 107.

¹H NMR (CDCl₃) δ: 1.77 (4H, s), 2.05 (3H, s), 2.51 (4H, s),
3.25 (2H, s), 4.74 (2H, s), 7.14-7.50 (6H, m), 7.63 (2H,
5 d, J=7.2 Hz), 7.71 (2H, d, J=8.4 Hz), 7.79 (1H, s), 7.94
(2H, d, J=8.4 Hz).

Melting point: 176-178 °C (crystallization solvent:
ethyl acetate - diisopropyl ether)

10 Example 214

4'-Methoxy-N-[4-methyl-3-(1-pyrrolidinymethyl)-2H-
chromen-7-yl][1,1'-biphenyl]-4-carboxamide



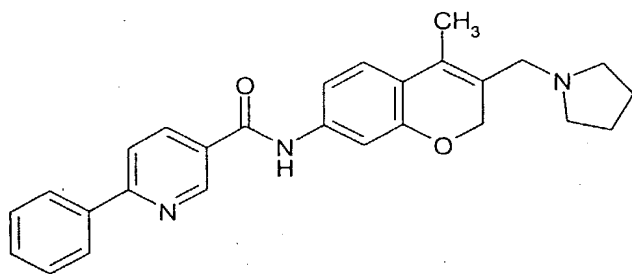
The titled compound was obtained as colorless powders
15 by carrying out the same operation as in Example 1, using
4-methyl-3-(1-pyrrolidinymethyl)-2H-chromen-7-amine
obtained in Reference Example 107.

¹H NMR (CDCl₃) δ: 1.77 (4H, s), 2.05 (3H, s), 2.51 (4H, s),
3.25 (2H, s), 3.87 (3H, s), 4.74 (2H, s), 7.01 (2H, d, J=8.7
20 Hz), 7.14-7.31 (3H, m), 7.57 (2H, d, J=8.7 Hz), 7.66 (2H,
d, J=8.4 Hz), 7.89 (1H, s), 7.91 (2H, d, J=8.4 Hz).

Melting point: 195-197 °C (crystallization solvent:
ethyl acetate - diisopropyl ether)

25 Example 215

N-[4-Methyl-3-(1-pyrrolidinymethyl)-2H-chromen-7-yl]-
6-phenylnicotinamide



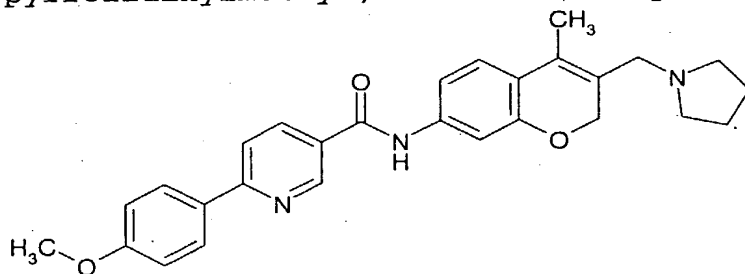
The titled compound was obtained as colorless powders by carrying out the same operation as in Example 1, using 4-methyl-3-(1-pyrrolidinylmethyl)-2H-chromen-7-amine obtained in Reference Example 107.

¹H NMR (CDCl₃) δ: 1.77 (4H, s), 2.05 (3H, s), 2.51 (4H, s), 3.25 (2H, s), 4.74 (2H, s), 7.14-7.28 (3H, m), 7.47-7.54 (3H, m), 7.81-7.87 (2H, m), 8.06 (2H, d, J=8.4 Hz), 8.27 (1H, d, J=8.4 Hz), 9.13 (1H, s).

Melting point: 192-193 °C (crystallization solvent: ethyl acetate)

Example 216

6-(4-Methoxyphenyl)-N-[4-methyl-3-(1-pyrrolidinylmethyl)-2H-chromen-7-yl]nicotinamide



The titled compound was obtained as colorless powders by carrying out the same operation as in Example 1, using 4-methyl-3-(1-pyrrolidinylmethyl)-2H-chromen-7-amine obtained in Reference Example 107.

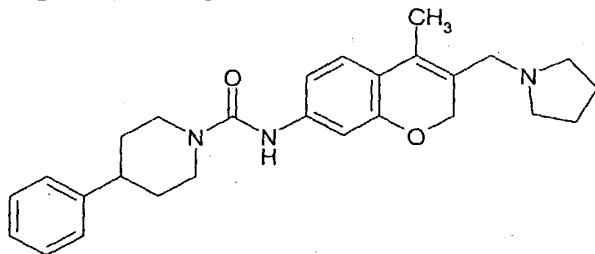
¹H NMR (CDCl₃) δ: 1.77 (4H, s), 2.05 (3H, s), 2.51 (4H, s), 3.25 (2H, s), 3.89 (3H, s), 4.74 (2H, s), 7.03 (2H, d, J=8.7 Hz), 7.14-7.26 (3H, m), 7.75-7.81 (2H, m), 8.03 (2H, d, J=8.7 Hz), 8.21 (1H, d, J=6.6 Hz), 9.09 (1H, s).

Melting point: 201-203 °C (crystallization solvent: ethyl

acetate)

Example 217

N-[4-Methyl-3-(1-pyrrolidinylmethyl)-2H-chromen-7-yl]-
5 4-phenyl-1-piperidinecarboxamide



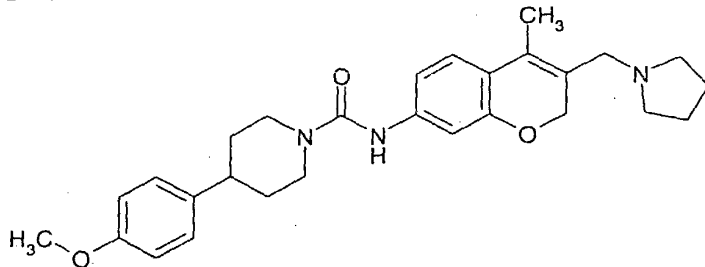
The titled compound was obtained as colorless powders
by carrying out the same operation as in Example 99, using
4-methyl-3-(1-pyrrolidinylmethyl)-2H-chromen-7-amine
10 obtained in Reference Example 107.

¹H NMR (CDCl₃) δ: 1.72-1.95 (8H, m), 2.03 (3H, s), 2.54 (4H,
s), 2.63-2.76 (1H, m), 2.95-3.00 (2H, m), 3.27 (2H, s),
4.19-4.23 (2H, m), 4.70 (2H, s), 6.39 (1H, s), 6.83 (1H,
s), 7.01-7.32 (7H, m).

15 Melting point: 125-127 °C (crystallization solvent:
ethyl acetate - diisopropyl ether)

Example 218

4-(4-Methoxyphenyl)-N-[4-methyl-3-(1-
20 pyrrolidinylmethyl)-2H-chromen-7-yl]-1-
piperidinecarboxamide



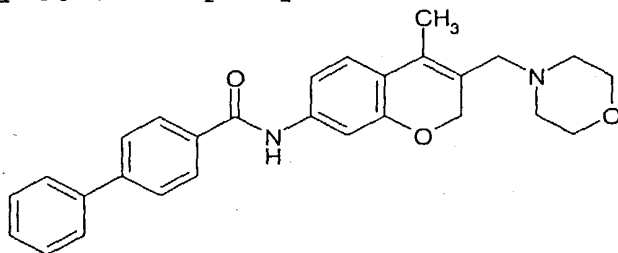
The titled compound was obtained as colorless powders
by carrying out the same operation as in Example 99, using
25 4-methyl-3-(1-pyrrolidinylmethyl)-2H-chromen-7-amine

obtained in Reference Example 107.

¹H NMR (CDCl₃) δ: 1.63-1.91 (8H, m), 2.02 (3H, s), 2.49 (4H, s), 2.61-2.71 (1H, m), 2.93-3.01 (2H, m), 3.23 (2H, s), 3.79 (3H, s), 4.16-4.21 (2H, m), 4.69 (2H, s), 6.34 (1H, s),
5 6.82-6.91 (3H, m), 6.99-7.02 (1H, m), 7.10-7.15 (3H, m).
Melting point: 144-146 °C (crystallization solvent: ethyl acetate - n-hexane)

Example 219

10 N-[4-Methyl-3-(4-morpholinylmethyl)-2H-chromen-7-yl][1,1'-biphenyl]-4-carboxamide



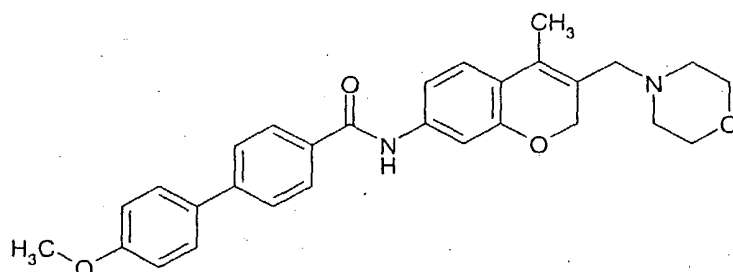
The titled compound was obtained as colorless powders by carrying out the same operation as in Example 1, using
15 4-methyl-3-(4-morpholinylmethyl)-2H-chromen-7-amine
obtained in Reference Example 108.

¹H NMR (DMSO-d₆) δ: 2.01 (3H, s), 2.37 (4H, s), 3.32 (2H, s), 3.57 (4H, s), 4.63 (2H, s), 7.23 (1H, d, J=8.1 Hz),
20 7.38-7.54 (5H, m), 7.76 (2H, d, J=7.5 Hz), 7.84 (2H, d, J=8.1 Hz), 8.04 (2H, d, J=8.1 Hz), 10.27 (1H, s).

Melting point: 162-164 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 220

25 4'-Methoxy-N-[4-methyl-3-(4-morpholinylmethyl)-2H-chromen-7-yl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained as colorless powders by carrying out the same operation as in Example 1, using 4-methyl-3-(4-morpholinylmethyl)-2H-chromen-7-amine

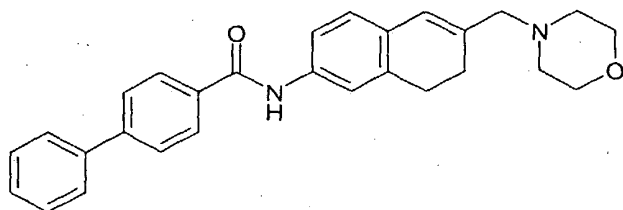
5 obtained in Reference Example 108.

¹H NMR (DMSO-d₆) δ: 2.00 (3H, s), 2.37 (4H, s), 3.11 (2H, s), 3.57 (4H, s), 3.82 (3H, s), 4.63 (2H, s), 7.07 (2H, d, J=8.7 Hz), 7.23 (1H, d, J=8.1 Hz), 7.38-7.40 (2H, m), 7.72 (2H, d, J=8.7 Hz), 7.79 (2H, d, J=8.4 Hz), 8.01 (2H, d, J=8.4 Hz), 10.23 (1H, s).

10 Melting point: 198-200 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 221

15 N-[6-(4-Morpholinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained as colorless powders by carrying out the same operation as in Example 1, using 6-(4-morpholinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 109.

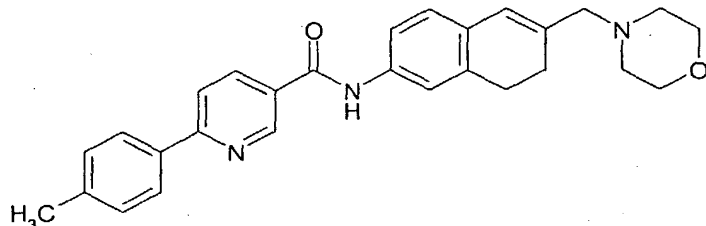
¹H-NMR (CDCl₃) δ: 2.34 (2H, t, J=8.4 Hz), 2.45 (4H, m), 2.85 (2H, t, J=8.4 Hz), 3.06 (2H, s), 3.73 (4H, t, J=4.7 Hz), 6.36 (1H, s), 7.02 (1H, d, J=8.1 Hz), 7.36-7.78 (10H, m), 7.93 (2H, d, J=8.1 Hz).

25 Melting point: 180-181 °C (crystallization solvent:

ethyl acetate - diisopropyl ether)

Example 222

6-(4-Methylphenyl)-N-[6-(4-morpholinylmethyl)-7,8-
5 dihydro-2-naphthalenyl]nicotinamide



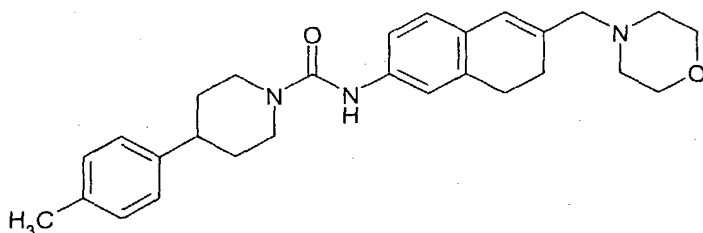
The titled compound was obtained as colorless powders by carrying out the same operation as in Example 1, using 6-(4-morpholinylmethyl)-7,8-dihydro-2-naphthalenamine
10 obtained in Reference Example 109.

¹H-NMR (CDCl₃) δ: 2.39 (2H, t, J=8.4 Hz), 2.43 (7H, m), 2.85 (2H, t, J=8.4 Hz), 3.06 (2H, s), 3.73 (4H, t, J=4.5 Hz), 6.36 (1H, s), 7.03 (1H, d, J=8.1 Hz), 7.30-7.38 (3H, m), 7.50 (1H, s), 7.76 (1H, s), 7.84 (1H, d, J=8.1 Hz), 7.97
15 (2H, d, J=8.1 Hz), 8.24 (1H, dd, J=8.4, 2.3 Hz), 9.12 (1H, s).

Melting point: 233-234 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

20 Example 223

4-(4-Methylphenyl)-N-[6-(4-morpholinylmethyl)-7,8-
dihydro-2-naphthalenyl]-1-piperidinecarboxamide



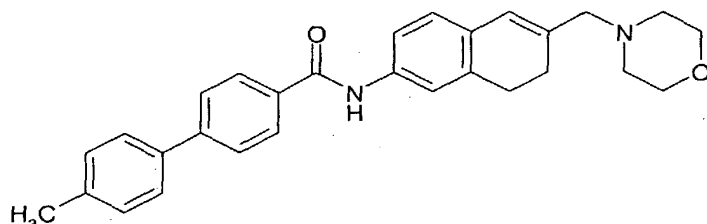
The titled compound was obtained as colorless powders by carrying out the same operation as in Example 99, using 6-(4-morpholinylmethyl)-7,8-dihydro-2-naphthalenamine
25 obtained in Reference Example 109.

¹H-NMR (CDCl₃) δ: 1.65-1.75 (4H, m), 1.90 (2H, m), 2.27-2.43 (7H, m), 2.72 (1H, m), 2.79 (2H, t, J=7.5 Hz), 2.93-3.04 (4H, m), 3.72 (4H, m), 4.20 (2H, d, J=11.7 Hz), 6.31 (1H, s), 6.39 (1H, s), 6.92 (1H, d, J=8.1 Hz), 7.05-7.26 (6H, m).

Melting point: 231-214 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 224

4'-Methyl-N-[6-(4-morpholinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



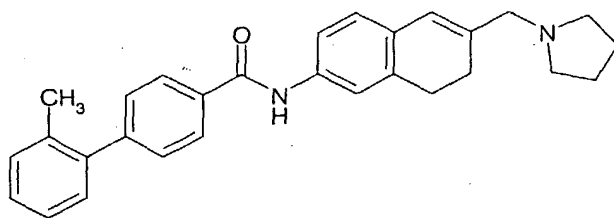
The titled compound was obtained as colorless powders by carrying out the same operation as in Example 1, using 6-(4-morpholinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 109.

¹H-NMR (CDCl₃) δ: 2.33 (2H, t, J=8.1 Hz), 2.42-2.44 (7H, m), 2.84 (2H, t, J=8.1 Hz), 3.06 (2H, s), 3.72 (4H, t, J=4.2 Hz), 6.36 (1H, s), 7.01 (1H, d, J=8.1 Hz), 7.25-7.29 (2H, m), 7.37 (1H, d, J=8.1 Hz), 7.51-7.54 (3H, m), 7.68 (2H, d, J=8.1 Hz), 7.85 (1H, s), 7.92 (2H, d, J=8.1 Hz).

Melting point: 196-197 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 225

2'-Methyl-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide

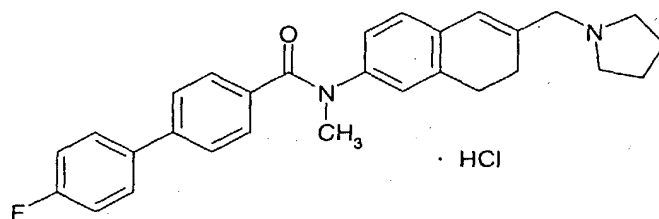


The titled compound was obtained as colorless powders by carrying out the same operation as in Example 1, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine
 5 obtained in Reference Example 54.

Melting point: 177-178 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 226

10 4'-Fluoro-N-methyl-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide Hydrochloride

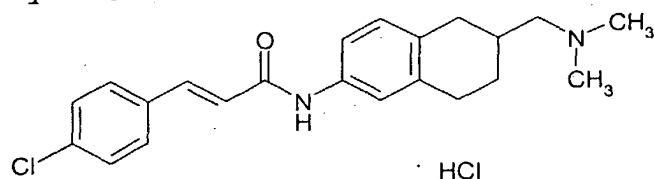


15 N-Methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine dihydrochloride (315 mg, 1.0 mmol) obtained in Reference Example 113 was dissolved in N,N-dimethylformamide (25 ml). 4-Bromobenzoic acid (402 mg, 2.0 mmol), WSC (383 mg, 2.0 mmol), HOBt (270 mg, 2.0 mmol) and DMAP (244mg, 2.0 mmol) were added to the solution,
 20 which was stirred at room temperature for 16 hours. Ethyl acetate and water were added to the reaction mixture, and extraction was conducted. The ethyl acetate layer was concentrated under reduced pressure. The residue was purified by aluminum column chromatography (development
 25 solvent; ethyl acetate : n-hexane = 33:67). The eluate was concentrated under reduced pressure, which was dissolved in dimethoxyethane - tetrahydrofuran (10:1, 5.5 ml).

4-Fluorophenylboric acid (73 mg, 0.52 mmol) ,
 tetrakis(triphenylphosphine)palladium complex (15 mg,
 0.013 mmol) and 2N aqueous sodium carbonate solution (0.433 ml) were added to the solution, which was refluxed
 5 with heating under nitrogen atmosphere at 90°C for 5.5 hours. The reaction mixture was poured into cold water, and extraction was conducted using ethyl acetate. The ethyl acetate layer was concentrated, and the residue was purified by aluminum column chromatography (development
 10 solvent; ethyl acetate) . 4N Hydrogen chloride - ethyl acetate solution was added to the eluate, which was concentrated under reduced pressure. The resulting residue was recrystallized from methanol - ethyl acetate, to give the titled compound (108 mg) .
 15 ¹H-NMR (DMSO-d₆) δ: 1.92-1.98 (4H, m), 2.39 (2H, t, J=8.1 Hz), 2.73 (2H, t, J=8.1 Hz), 3.00 (2H, m), 3.35 (3H, m), 3.44 (2H, m), 3.83 (2H, d, J=5.6 Hz), 6.62 (1H, s), 6.92-7.01 (2H, m), 7.11 (1H, s), 7.26 (2H, dd, J=8.9, 5.6 Hz), 7.38 (2H, d, J=8.1 Hz), 7.55 (2H, d, J=8.1 Hz), 7.69 (2H, dd,
 20 J=8.9, 5.6 Hz), 10.60 (1H, brs.).
 Melting point: 201-203 °C (crystallization solvent: methanol - diisopropyl ether)
 FAB(pos) 441.2 [M+H]⁺

25 Example 227

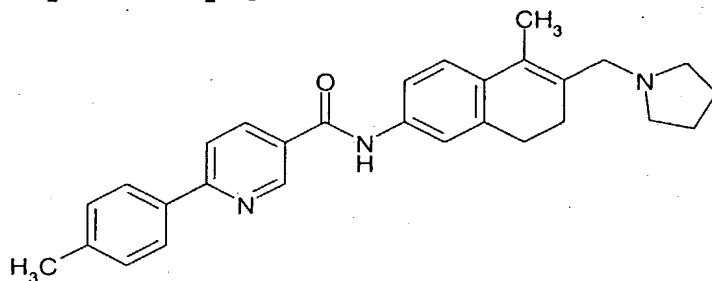
(E)-3-(4-Chlorophenyl)-N-[6-[(dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-2-propenamide
 Hydrochloride



30 The titled compound was obtained as colorless powders by carrying out the same operation as in Example 4.
 Melting point: 243-245 °C (crystallization solvent: methanol - diisopropyl ether)

Example 228

6-(4-Methylphenyl)-N-[5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]nicotinamide



The titled compound was obtained as colorless powders by carrying out the same operation as in Example 1, using 5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 69.

Melting point: 175-176 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

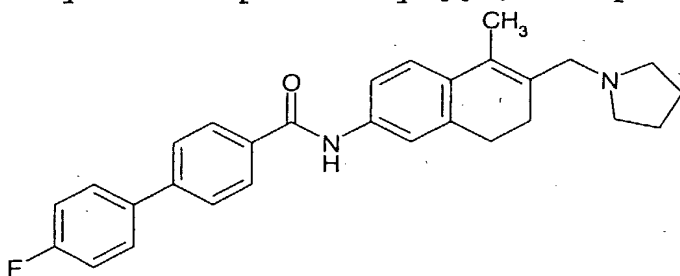
Elemental analysis for $C_{29}H_{30}N_3O$

Calcd.: C, 79.78; H, 6.93; N, 9.63

Found: C, 79.66; H, 6.97; N, 9.68

Example 229

4'-Fluoro-N-[5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained as colorless powders by carrying out the same operation as in Example 1, using 5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 69.

Melting point: 199-201 °C (crystallization solvent: ethyl

acetate - diisopropyl ether)

Elemental analysis for $C_{29}H_{30}FN_2O$

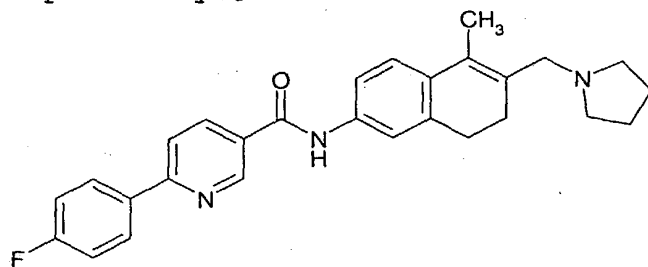
Calcd.: C, 79.06; H, 6.63; N, 6.36

Found: C, 79.01; H, 6.81; N, 6.45

5

Example 230

6-(4-Fluorophenyl)-N-[5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]nicotinamide



10

The titled compound was obtained as colorless powders by carrying out the same operation as in Example 1, using 5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 69.

15 Melting point: 204-205 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Elemental analysis for $C_{28}H_{28}FN_3O$

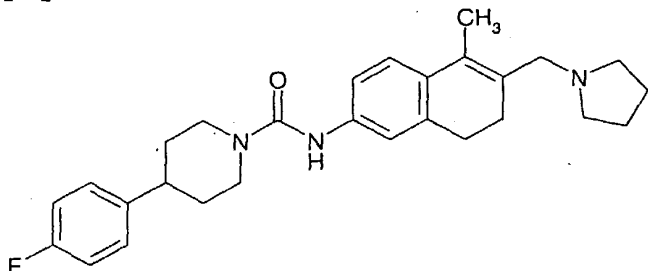
Calcd.: C, 76.17; H, 6.39; N, 9.52

Found: C, 76.03; H, 6.44; N, 9.62

20

Example 231

4-(4-Fluorophenyl)-N-[5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide



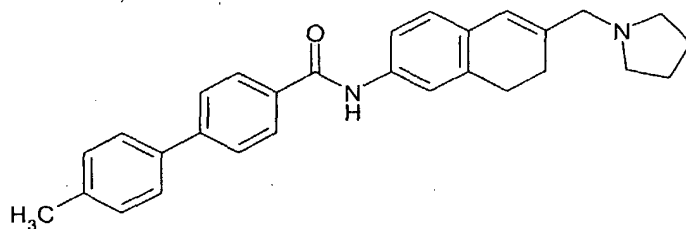
25

The titled compound was obtained as colorless powders by carrying out the same operation as in Example 99, using 5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 69.

5 Melting point: 172-173 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 232

4'-Methyl-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide

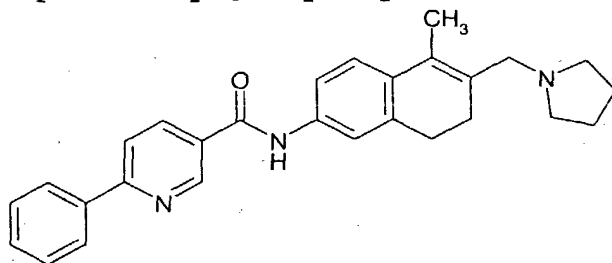


The titled compound was obtained as colorless powders by carrying out the same operation as in Example 1, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

15 Melting point: 176-177 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 233

20 N-[5-Methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]-6-phenylnicotinamide



The titled compound was obtained as colorless powders by carrying out the same operation as in Example 1, using 5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 69.

Melting point: 178-179 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

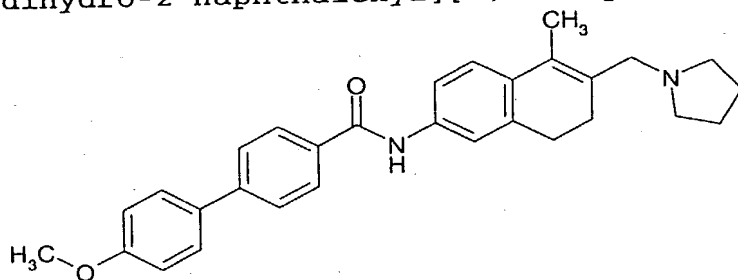
Elemental analysis for $C_{28}H_{29}N_3O$

Calcd.: C, 79.40; H, 6.90; N, 9.92

5 Found: C, 79.13; H, 6.82; N, 10.03

Example 234

4'-Methoxy-N-[5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



10

The titled compound was obtained as colorless powders by carrying out the same operation as in Example 1, using 5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 69.

15 1H -NMR ($CDCl_3$) δ : 1.78 (4H, m), 2.10 (3H, s), 2.37 (2H, t, $J=8.1$ Hz), 2.53 (4H, m), 2.76 (2H, t, $J=8.1$ Hz), 3.28 (2H, s), 3.87 (3H, s), 7.01 (1H, d, $J=8.6$ Hz), 7.27 (2H, d, $J=7.8$ Hz), 7.46 (1H, d, $J=7.8$ Hz), 7.48 (1H, s), 7.57 (2H, d, $J=8.6$ Hz), 7.66 (2H, d, $J=8.6$ Hz), 7.81 (1H, s), 7.92 (2H, d, $J=7.8$ Hz).

20

Melting point: 179-180 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Elemental analysis for $C_{30}H_{32}N_2O_2$

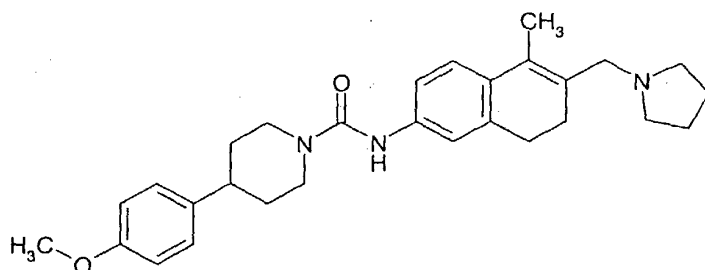
Calcd.: C, 79.61; H, 7.13; N, 6.19

25 Found: C, 79.35; H, 7.28; N, 6.24

Example 235

4-(4-Methoxyphenyl)-N-[5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide

30



The titled compound was obtained as colorless powders by carrying out the same operation as in Example 99, using 5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 69.

¹H-NMR (CDCl₃) δ: 1.67 (2H, dd, J=13.4, 4.0 Hz), 1.78 (4H, m), 1.89 (2H, d, J=11.4 Hz), 2.07 (3H, s), 2.34 (2H, t, J=7.5 Hz), 2.52 (4H, m), 2.68-2.73 (3H, m), 2.98 (2H, t, J=7.5 Hz), 3.26 (2H, s), 3.80 (3H, s), 4.20 (2H, d, J=13.4 Hz), 6.36 (1H, s), 6.86 (2H, d, J=8.4 Hz), 7.12-7.20 (5H, m).

Melting point: 163-164 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

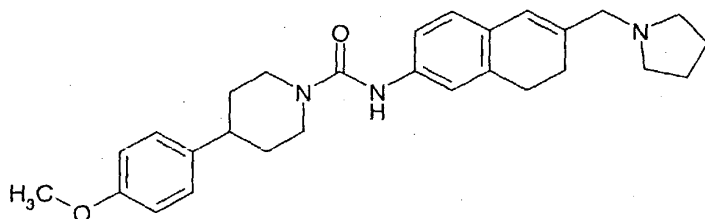
Elemental analysis for C₂₈H₃₇N₃O₂

Calcd.: C, 75.13; H, 8.33; N, 9.39

Found: C, 74.96; H, 8.14; N, 9.10

Example 236

4-(4-Methoxyphenyl)-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide



The titled compound was obtained as colorless powders by carrying out the same operation as in Example 1, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

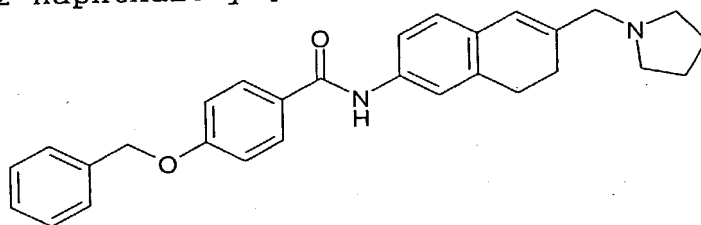
¹H-NMR (CDCl₃) δ: 1.61-1.91 (8H, m), 2.31 (2H, t, J=8.1 Hz), 2.54 (4H, m), 2.73-2.81 (3H, m), 2.98 (2H, t, J=7.8 Hz),

3.16 (2H, s), 3.79 (3H, s), 4.20 (2H, d, $J=13.1$ Hz), 6.31 (1H, s), 6.36 (1H, s), 6.86 (2H, d, $J=8.6$ Hz), 7.06-7.20 (5H, m).

Melting point: 175-176 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 237

4-(Benzyloxy)-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]benzamide



The titled compound was obtained as colorless powders by carrying out the same operation as in Example 1, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

Melting point: 174-175 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

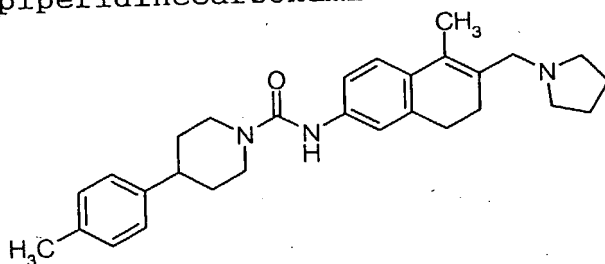
Elemental analysis for $C_{28}H_{30}N_2O_2$

Calcd.: C, 78.84; H, 7.09; N, 6.87

Found: C, 79.06; H, 6.99; N, 6.41

Example 238

4-(4-Methylphenyl)-N-[5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide



The titled compound was obtained by carrying out the same operation as in Example 99, using 5-methyl-6-(1-

pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine
obtained in Reference Example 69.

¹H-NMR (CDCl₃) δ: 1.65-1.78 (6H, m), 1.90 (2H, d, J=12.9 Hz), 2.07 (3H, s), 2.33-2.37 (5H, m), 2.53 (4H, m),
5 2.68-2.74 (3H, m), 2.99 (2H, m), 3.27(2H,s), 4.21 (2H, d, J=13.2 Hz), 6.37 (1H, s), 7.09-7.21 (7H, m).

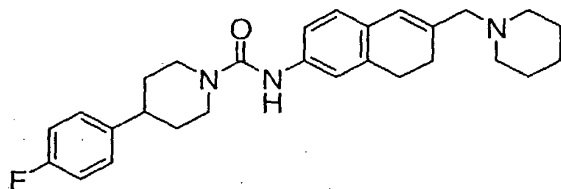
Melting point: 159-160 °C (crystallization solvent:
ethyl acetate - diisopropyl ether)

FAB(pos) 444.3 [M+H]⁺

10

Example 239

4-(4-Fluorophenyl)-N-[6-(1-piperidinylmethyl)-7,8-
dihydro-2-naphthalenyl]-1-piperidinecarboxamide



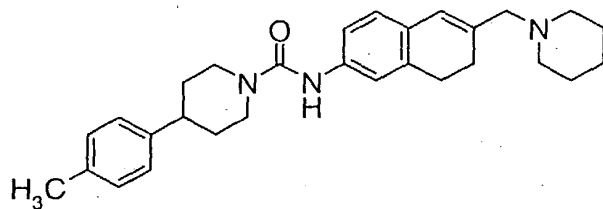
15 The titled compound was obtained by carrying out the
same operation as in Example 99, using 6-(1-
piperidinylmethyl)-7,8-dihydro-2-naphthalenamine
dihydrochloride obtained in Reference Example 114.

¹H-NMR (CDCl₃) δ: 1.43 (2H, m), 1.56-1.75 (6H, m), 1.89 (2H,
20 d, J=12.3 Hz), 2.27-2.36 (6H, m), 2.70 (1H, m), 2.78 (2H,
t, J=7.5 Hz), 2.88-3.00 (4H, m), 4.20 (2H, d, J=13.2 Hz),
6.29 (1H, s), 6.38 (1H, s), 6.91-7.08 (4H, m), 7.14-7.20 (3H,
m).

Melting point: 194 -195 °C (crystallization solvent:
25 ethyl acetate - diisopropyl ether)

Example 240

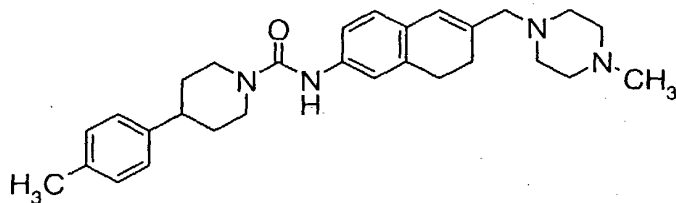
4-(4-Methylphenyl)-N-[6-(1-piperidinylmethyl)-7,8-
dihydro-2-naphthalenyl]-1-piperidinecarboxamide



- The titled compound was obtained by carrying out the same operation as in Example 99, using 6-(1-piperidinylmethyl)-7,8-dihydro-2-naphthalenamine dihydrochloride obtained in Reference Example 114.
- ¹H-NMR (CDCl₃) δ: 1.43 (2H, m), 1.56-1.74 (6H, m), 1.90 (2H, d, J=12.0 Hz), 2.27-2.36 (9H, m), 2.69 (1H, m), 2.79 (2H, t, J=8.1 Hz), 2.94-3.01 (4H, m), 4.19 (2H, d, J=13.2 Hz), 6.29 (1H, s), 6.35 (1H, s), 6.93 (2H, d, J=8.1 Hz), 7.05-7.26 (5H, m).
- Melting point: 209 -210 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 241

- 4-(4-Methylphenyl)-N-[6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide



- The titled compound was obtained by carrying out the same operation as in Example 99, using 6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenamine obtained in Reference Example 106.
- ¹H NMR (CDCl₃) δ: 1.62-1.77 (2H, m), 1.90 (2H, d, J=12.0 Hz), 2.28 (2H, t, J=8.1 Hz), 2.29 (3H, s), 2.33 (3H, s), 2.46 (8H, bs), 2.64-2.73 (1H, m), 2.79 (2H, t, J=8.1 Hz), 2.96 (2H, d, J=10.5 Hz), 3.05 (2H, s), 4.19 (2H, d, J=13.5 Hz), 6.31 (1H, s), 6.34 (1H, s), 6.93 (1H, d, J=8.4 Hz), 7.04-7.16 (5H, m), 7.23 (1H, s).

Melting point: 214-216 °C (crystallization solvent: tetrahydrofuran - n-hexane)

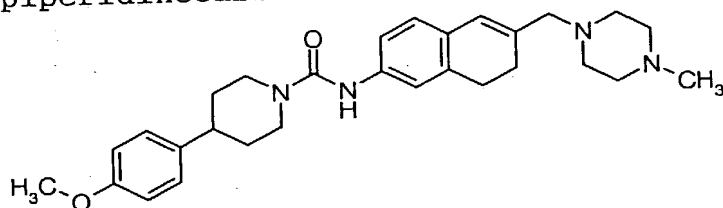
Elemental analysis for $C_{29}H_{38}N_4O$

Calcd.: C, 75.94; H, 8.35; N, 12.22.

5 Found: C, 75.67; H, 8.47; N, 12.27.

Example 242

4-(4-Methoxyphenyl)-N-[6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide



The titled compound was obtained by carrying out the same operation as in Example 99, using 6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenamine

15 obtained in Reference Example 106.
 1H NMR ($CDCl_3$) δ : 1.68-1.76 (2H, m), 1.89 (2H, d, $J=11.1$ Hz), 2.29 (2H, t, $J=8.1$ Hz), 2.29 (3H, s), 2.46 (8H, bs), 2.64-2.71 (1H, m), 2.79 (2H, t, $J=8.1$ Hz), 2.82-3.03 (2H, m), 3.05 (2H, s), 3.80 (3H, s), 4.19 (2H, d, $J=12.6$ Hz),
 20 6.31 (1H, s), 6.34 (1H, s), 6.87 (2H, d, $J=8.7$ Hz), 6.93 (1H, d, $J=8.4$ Hz), 7.06 (1H, dd, $J=8.1, 2.1$ Hz), 7.14 (2H, d, $J=8.7$ Hz), 7.23 (1H, s).

Melting point: 198-200 °C (crystallization solvent: tetrahydrofuran - n-hexane)

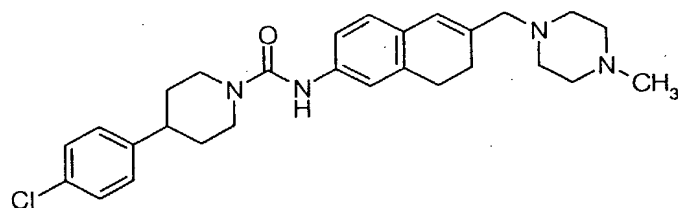
25 Elemental analysis for $C_{29}H_{38}N_4O_2$

Calcd.: C, 73.38; H, 8.07; N, 11.80.

Found: C, 73.04; H, 7.95; N, 11.67.

Example 243

30 4-(4-Chlorophenyl)-N-[6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide



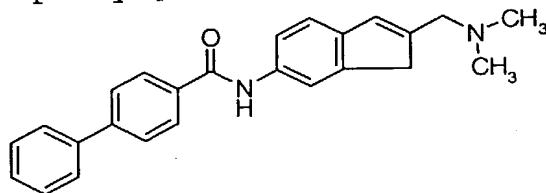
The titled compound was obtained by carrying out the same operation as in Example 99, using 6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenamine obtained in Reference Example 106.

¹H NMR (CDCl₃) δ: 1.64-1.76 (2H, m), 1.90 (2H, d, J=11.1 Hz), 2.29 (2H, t, J=8.1 Hz), 2.29 (3H, s), 2.46 (8H, bs), 2.66-2.72 (1H, m), 2.79 (2H, t, J=8.1 Hz), 2.81-3.03 (2H, m), 3.05 (2H, s), 4.20 (2H, d, J=12.6 Hz), 6.31 (1H, s), 6.34 (1H, s), 6.93 (1H, d, J=7.8 Hz), 7.04-7.07 (1H, m), 7.14 (2H, d, J=8.4 Hz), 7.22 (1H, s), 7.28 (2H, d, J=8.4 Hz).

Melting point: 201-203 °C (crystallization solvent: tetrahydrofuran - n-hexane)

Example 244

N-[2-[(Dimethylamino)methyl]-1H-inden-6-yl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 2-[(dimethylamino)methyl]-1H-inden-6-amine obtained in Reference Example 116.

Elemental analysis for C₂₅H₂₄N₂O · 0.5H₂O

Calcd.: C, 79.55; H, 6.68; N, 7.42.

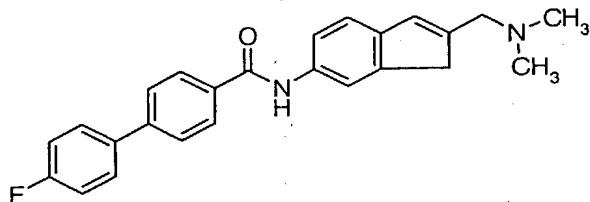
Found: C, 79.38; H, 6.76; N, 7.34.

Melting point: 187-189 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

FAB(pos) 369.2 [M+H]⁺

Example 245

N-[2-[(Dimethylamino)methyl]-1H-inden-6-yl]-4'-
5 fluoro[1,1'-biphenyl]-4-carboxamide



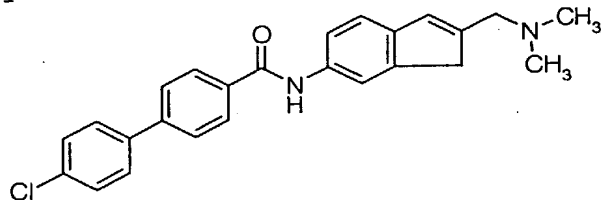
The titled compound was obtained by carrying out the
same operation as in Example 1, using 2-
[(dimethylamino)methyl]-1H-inden-6-amine obtained in
10 Reference Example 116.

Melting point: 209-211 °C (crystallization solvent: ethyl
acetate - diisopropyl ether)

FAB(pos) 387.2 [M+H]⁺

15 Example 246

4'-Chloro-N-[2-[(dimethylamino)methyl]-1H-inden-6-
yl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the
20 same operation as in Example 1, using 2-
[(dimethylamino)methyl]-1H-inden-6-amine obtained in
Reference Example 116.

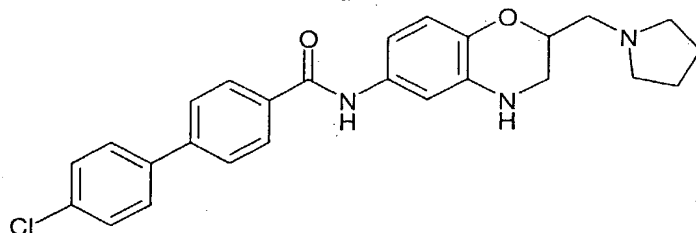
Melting point: 218-220 °C (crystallization solvent: ethyl
acetate - diisopropyl ether)

25 FAB(pos) 403.2 [M+H]⁺

Example 247

4'-Chloro-N-[2-(1-pyrrolidinylmethyl)-3,4-dihydro-2H-

1,4-benzoxazin-6-yl][1,1'-biphenyl]-4-carboxamide



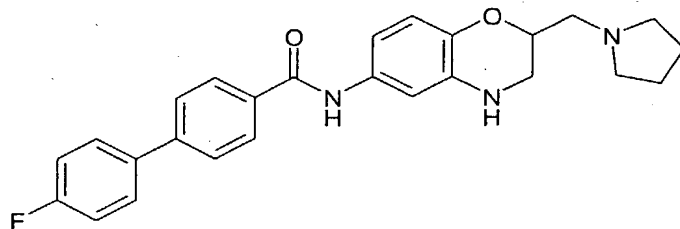
The titled compound was obtained by carrying out the same operation as in Example 1, using 6-amino-2-(1-pyrrolidinylmethyl)-3,4-dihydro-2H-1,4-benzoxazine obtained in Reference Example 117.

¹H-NMR (CDCl₃) δ: 1.70-1.90 (4H, m), 2.50-2.70 (4H, m), 2.73 (2H, d, J=6.0Hz), 3.18-3.24 (1H, m), 3.45-3.49 (1H, m), 3.87 (1H, brs), 4.26-4.28 (1H, m), 6.61 (1H, dd, J=2.7, 8.4 Hz), 6.80 (1H, d, J=8.4 Hz), 7.26 (1H, d, J=2.7 Hz), 7.44 (2H, d, J=8.4 Hz), 7.55 (2H, d, J=8.4 Hz), 7.64 (2H, d, J=8.1 Hz), 7.71 (1H, s), 7.91 (2H, d, J=8.1 Hz).

Melting point: 221-222 °C (crystallization solvent: diisopropyl ether)

Example 248

4'-Fluoro-N-[2-(1-pyrrolidinylmethyl)-3,4-dihydro-2H-1,4-benzoxazin-6-yl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the same operation as in Reference Example 1, using 6-amino-2-(1-pyrrolidinylmethyl)-3,4-dihydro-2H-1,4-benzoxazine obtained in Reference Example 117.

¹H-NMR (CDCl₃) δ: 1.70-1.90 (4H, m), 2.50-2.70 (4H, m), 2.73 (2H, d, J = 6.3Hz), 3.18-3.24 (1H, m), 3.45-3.49 (1H, m), 3.88 (1H, brs), 4.24-4.30 (1H, m), 6.62 (1H, dd, J=2.7, 8.4 Hz), 6.80 (1H, d, J=8.4 Hz), 7.13-7.19 (2H, m), 7.26 (1H,

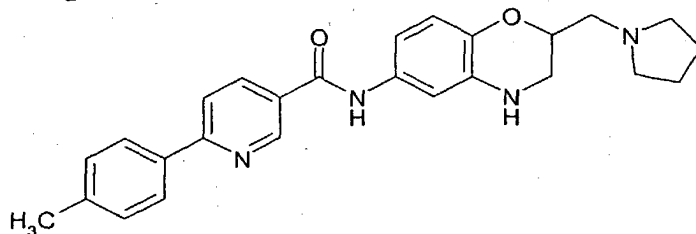
d, $J=2.7$ Hz), 7.56-7.60 (2H, m), 7.63 (2H, d, $J=8.4$ Hz), 7.71 (1H, s), 7.90 (2H, d, $J=8.4$ Hz).

Melting point: 204-206 °C (crystallization solvent: diisopropyl ether)

5

Example 249

6-(4-Methylphenyl)-N-[2-(1-pyrrolidinylmethyl)-3,4-dihydro-2H-1,4-benzoxazin-6-yl]nicotinamide



10

The titled compound was obtained by carrying out the same operation as in Reference Example 1, using 6-amino-2-(1-pyrrolidinylmethyl)-3,4-dihydro-2H-1,4-benzoxazine obtained in Reference Example 117.

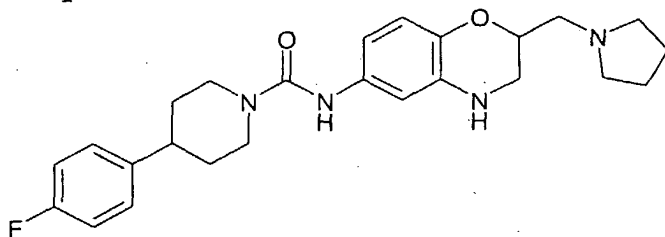
¹H-NMR (CDCl₃) δ : 1.70-1.85 (4H, m), 2.43 (3H, s), 2.50-2.70 (4H, m), 2.74 (2H, d, $J=6.3$ Hz), 3.19-3.25 (1H, m), 3.45-3.49 (1H, m), 3.90 (1H, brs), 4.27-4.29 (1H, m), 6.63 (1H, dd, $J=2.4, 8.7$ Hz), 6.81 (1H, d, $J=8.7$ Hz), 7.26 (1H, d, $J=2.7$ Hz), 7.31 (2H, d, $J=8.1$ Hz), 7.67 (1H, s), 7.81 (1H, d, $J=8.1$ Hz), 7.93 (2H, d, $J=7.8$ Hz), 8.21 (1H, dd, $J=2.4, 8.4$ Hz), 9.09 (1H, d, $J=2.4$ Hz).

20

Melting point: 207-208 °C (crystallization solvent: diisopropyl ether)

Example 250

4-(4-Fluorophenyl)-N-[2-(1-pyrrolidinylmethyl)-3,4-dihydro-2H-1,4-benzoxazin-6-yl]-1-piperidinecarboxamide



25

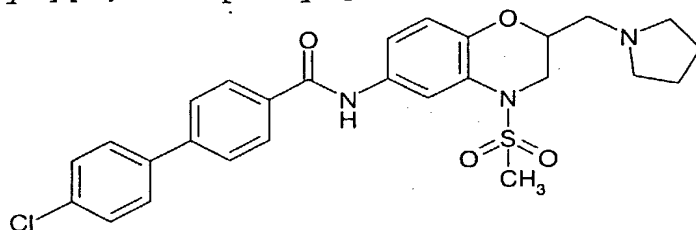
The titled compound was obtained by carrying out the same operation as in Reference Example 1, using 6-amino-2-(1-pyrrolidinylmethyl)-3,4-dihydro-2H-1,4-benzoxazine obtained in Reference Example 117.

5 ¹H-NMR (CDCl₃) δ : 1.60-1.90 (8H, m), 2.50-2.70 (5H, m), 2.71 (2H, d, J=6.3Hz), 2.91-3.00 (2H, m), 3.15-3.21 (1H, brs), 3.42-3.45 (1H, m), 3.77 (1H, brs), 4.15-4.25 (3H, m), 6.20 (1H, s), 6.38 (1H, dd, J=2.1, 8.4 Hz), 6.73 (1H, d, J=8.4 Hz), 6.91 (1H, d, J=2.1 Hz), 6.97-7.03 (2H, m), 7.14-7.19 (2H, m).

Melting point: 192-195 °C (crystallization solvent: diisopropyl ether)

Example 251

15 4'-Chloro-N-[4-(methanesulfonyl)-2-(1-pyrrolidinylmethyl)-3,4-dihydro-2H-1,4-benzoxazin-6-yl][1,1'-biphenyl]-4-carboxamide



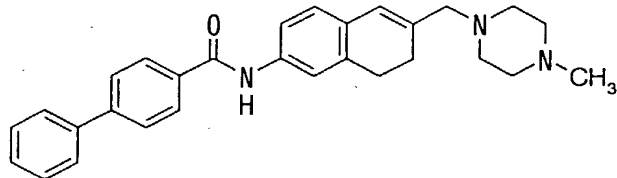
20 The titled compound was obtained by carrying out the same operation as in Reference Example 1, using 6-amino-4-(methanesulfonyl)-2-(1-pyrrolidinylmethyl)-3,4-dihydro-2H-1,4-benzoxazine obtained in Reference Example 118.

25 ¹H-NMR (CDCl₃) δ : 1.75-1.85 (4H, m), 2.55-2.70 (4H, m), 2.78 (2H, d, J=6.0Hz), 3.04 (3H, s), 3.27-3.34 (1H, m), 4.24-4.31 (1H, m), 4.31-4.35 (1H, m), 6.98 (1H, d, J=8.7 Hz), 7.45 (2H, d, J=9.0 Hz), 7.50-7.60 (1H, m), 7.53 (2H, d, J=9.0 Hz), 7.67 (2H, d, J=8.4 Hz), 7.84 (1H, s), 7.84 (1H, brs), 7.94 (2H, d, J=8.4 Hz).

30 Melting point: 203-204 °C (crystallization solvent: diisopropyl ether)

Example 252

N-[6-[(4-Methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



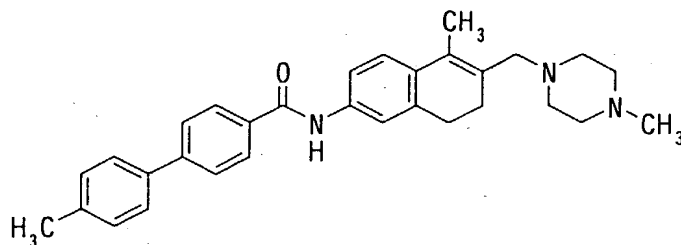
5 The titled compound was obtained by carrying out the same operation as in Reference Example 1, using 6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenamine obtained in Reference Example 106.

¹H NMR (CDCl₃) δ: 2.31 (3H, s), 2.33 (2H, t, J=8.1 Hz), 2.49
10 (8H, bs), 2.84 (2H, t, J=8.1 Hz), 3.07 (2H, s), 6.36 (1H, s), 7.02 (1H, d, J=8.1 Hz), 7.35-7.52 (5H, m), 7.63 (2H, d, J=8.1 Hz), 7.71 (2H, d, J=8.1 Hz), 7.80 (1H, s), 7.94 (2H, d, J=8.1 Hz).

Melting point: 196-198 °C (crystallization solvent:
15 ethyl acetate)

Example 253

4'-Methyl-N-[5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-
20 biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the same operation as in Reference Example 1, using 5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenamine obtained in Reference Example 115.

¹H NMR (CDCl₃) δ: 2.08 (3H, s), 2.29 (3H, s), 2.34 (2H, t, J=7.8 Hz), 2.42 (3H, s), 2.45 (8H, bs), 2.75 (2H, t, J=7.8 Hz), 3.16 (2H, s), 7.26-7.30 (3H, m), 7.44 (1H, d, J=8.4 Hz).

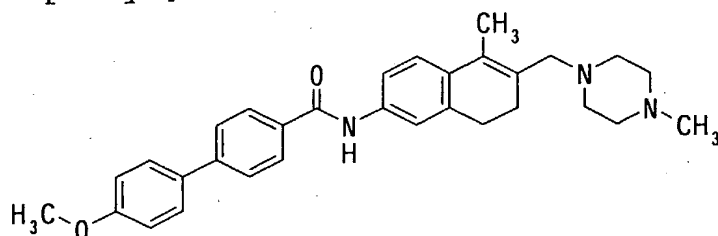
Hz), 7.53-7.55 (3H, m), 7.70 (2H, d, J=8.4 Hz), 8.00 (1H, s), 7.93 (2H, d, J=8.4 Hz).

Melting point: 212-214 °C (crystallization solvent: ethyl acetate)

5

Example 254

4'-Methoxy-N-[5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



10

The titled compound was obtained by carrying out the same operation as in Reference Example 1, using 5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenamine obtained in Reference Example 115.

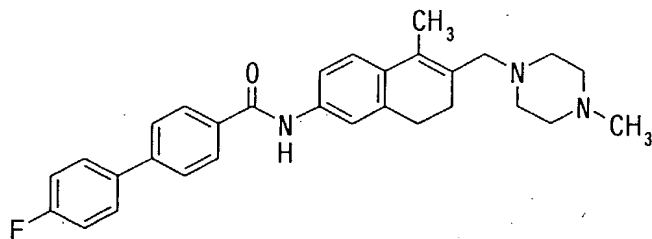
15 ¹H NMR (CDCl₃) δ: 2.08 (3H, s), 2.29 (3H, s), 2.34 (2H, t, J=7.8 Hz), 2.45 (8H, bs), 2.75 (2H, t, J=7.8 Hz), 3.16 (2H, s), 3.87 (3H, s), 7.01 (2H, d, J=8.1 Hz), 7.27 (1H, d, J=8.4 Hz), 7.44 (1H, d, J=8.4 Hz), 7.51 (1H, s), 7.58 (2H, d, J=8.4 Hz), 7.67 (2H, d, J=8.4 Hz), 7.81 (1H, s), 7.92 (2H, d, J=8.4 Hz).

20

Melting point: 215-217 °C (crystallization solvent: ethyl acetate)

Example 255

25 4'-Fluoro-N-[5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide

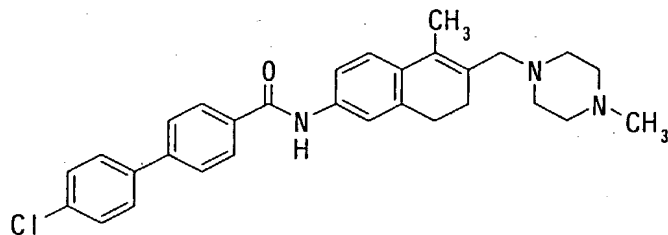


The titled compound was obtained by carrying out the same operation as in Reference Example 1, using 5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenamine obtained in Reference Example 115.

¹H NMR (CDCl₃) δ: 2.08 (3H, s), 2.29 (3H, s), 2.34 (2H, t, J=7.8 Hz), 2.46 (8H, bs), 2.75 (2H, t, J=7.8 Hz), 3.16 (2H, s), 7.17 (2H, d, J=8.4 Hz), 7.28 (1H, d, J=8.4 Hz), 7.44 (1H, d, J=8.4 Hz), 7.51 (1H, s), 7.57-7.62 (2H, m), 7.66 (2H, d, J=8.4 Hz), 7.82 (1H, s), 7.94 (2H, d, J=8.4 Hz).
Melting point: 233-235 °C (crystallization solvent: ethyl acetate)

Example 256

4'-Chloro-N-[5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



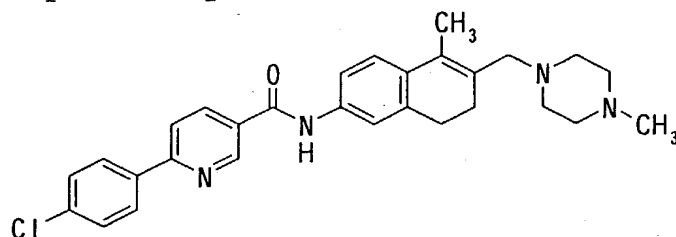
The titled compound was obtained by carrying out the same operation as in Reference Example 1, using 5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenamine obtained in Reference Example 115.

¹H NMR (CDCl₃) δ: 2.08 (3H, s), 2.29 (3H, s), 2.34 (2H, t, J=7.8 Hz), 2.46 (8H, bs), 2.75 (2H, t, J=7.8 Hz), 3.16 (2H, s), 7.28 (1H, d, J=8.4 Hz), 7.43-7.47 (3H, m), 7.51 (1H, s), 7.56 (2H, d, J=8.4 Hz), 7.67 (2H, d, J=8.4 Hz), 7.80 (1H, s), 7.94 (2H, d, J=8.4 Hz).

Melting point: 216-218 °C (crystallization solvent: ethyl acetate)

Example 257

5 6-(4-Chlorophenyl)-N-[5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl]nicotinamide



10 The titled compound was obtained by carrying out the same operation as in Reference Example 1, using 5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenamine obtained in Reference Example 115.

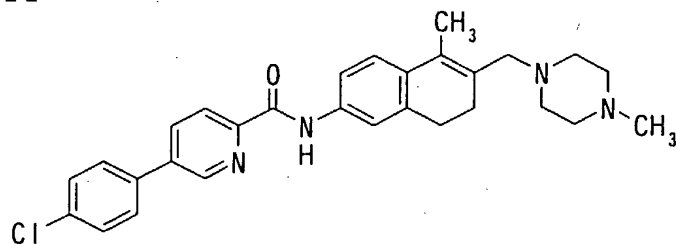
¹H NMR (CDCl₃) δ: 2.09 (3H, s), 2.29 (3H, s), 2.35 (2H, t, J=8.1 Hz), 2.46 (8H, bs), 2.75 (2H, t, J=8.1 Hz), 3.16 (2H, s), 7.28 (1H, d, J=8.4 Hz), 7.43-7.50 (4H, m), 7.83 (2H, d, J=8.4 Hz), 8.01 (2H, d, J=8.4 Hz), 8.27 (1H, d, J=8.4 Hz), 9.13 (1H, s).

15 Melting point: 219-221 °C (crystallization solvent: ethyl acetate)

20

Example 258

5-(4-Chlorophenyl)-N-[5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl]-2-pyridinecarboxamide



25

The titled compound was obtained by carrying out the same operation as in Reference Example 1, using 5-

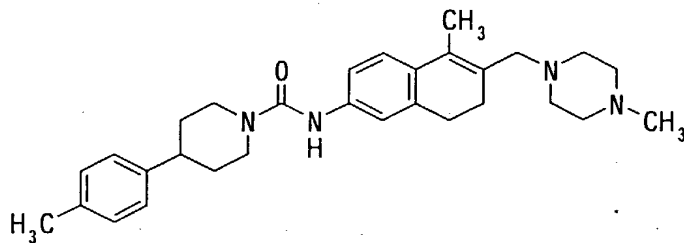
methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenamine obtained in Reference Example 115.

¹H NMR (CDCl₃) δ: 2.09 (3H, s), 2.29 (3H, s), 2.35 (2H, t, J=8.1 Hz), 2.45 (8H, bs), 2.77 (2H, t, J=8.1 Hz), 3.16 (2H, s), 7.30 (1H, d, J=8.1 Hz), 7.49-7.63 (6H, m), 8.05 (1H, dd, J=2.4 Hz, 8.4 Hz), 8.36 (1H, d, J=8.1 Hz), 8.79 (1H, d, J=1.2 Hz), 9.97 (1H, s).

Melting point: 177-179 °C (crystallization solvent: ethyl acetate)

Example 259

N-[5-Methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl]-4-(4-methylphenyl)-1-piperidinecarboxamide



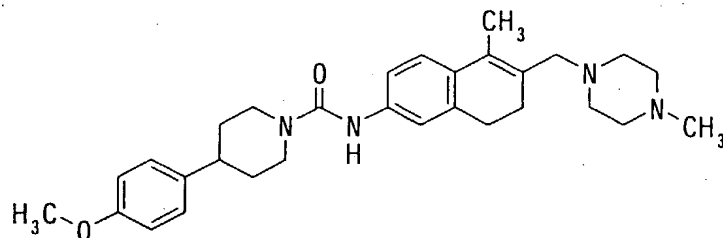
The titled compound was obtained by carrying out the same operation as in Reference Example 99, using 5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenamine obtained in Reference Example 115.

¹H NMR (CDCl₃) δ: 1.60-1.78 (4H, m), 2.05 (3H, s), 2.28 (3H, s), 2.29 (2H, t, J=8.1 Hz), 2.33 (3H, s), 2.46 (8H, bs), 2.65-2.72 (3H, m), 2.93-3.03 (2H, m), 3.13 (2H, s), 4.18-4.23 (2H, m), 6.40 (1H, s), 7.09-7.24 (7H, m).

Melting point: 176-178 °C (crystallization solvent: ethyl acetate-hexane)

Example 260

4-(4-Methoxyphenyl)-N-[5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide



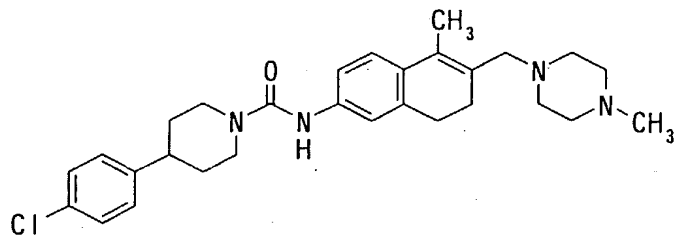
The titled compound was obtained by carrying out the same operation as in Reference Example 99, using 5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenamine obtained in Reference Example 115.

¹H NMR (CDCl₃) δ : 1.68-1.92 (4H, m), 2.05 (3H, s), 2.28 (3H, s), 2.29 (2H, t, J=8.1 Hz), 2.45 (8H, bs), 2.67-2.72 (3H, m), 2.95-3.02 (2H, m), 3.14 (2H, s), 3.80 (3H, s), 4.18-4.22 (2H, m), 6.36 (1H, s), 6.87 (2H, d, J=8.4 Hz), 7.12-7.21 (5H, m).

Melting point: 175-177 °C (crystallization solvent: ethyl acetate)

Example 261

4-(4-Chlorophenyl)-N-[5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide



The titled compound was obtained by carrying out the same operation as in Reference Example 99, using 5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenamine obtained in Reference Example 115.

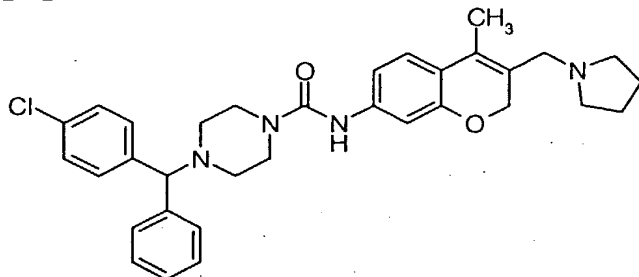
¹H NMR (CDCl₃) δ : 1.67-1.92 (4H, m), 2.05 (3H, s), 2.28 (3H, s), 2.29 (2H, t, J=8.1 Hz), 2.45 (8H, bs), 2.67-2.72 (3H, m), 2.95-3.02 (2H, m), 3.14 (2H, s), 4.18-4.23 (2H, m), 6.36 (1H, s), 7.13-7.30 (7H, m).

Melting point: 141-143 °C (crystallization solvent: ethyl acetate)

ethyl acetate)

Example 262

4-[(4-Chlorophenyl)(phenyl)methyl]-N-[4-methyl-3-(1-
pyrrolidinylmethyl)-2H-chromen-7-yl]-1-
piperazinecarboxamide



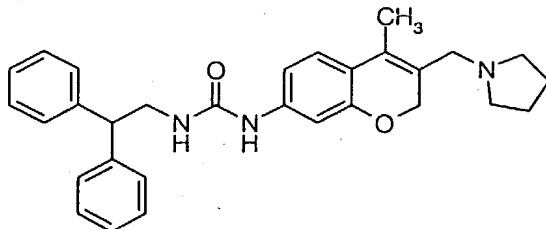
The titled compound was obtained by carrying out the
same operation as in Reference Example 99, using 4-
methyl-3-(1-pyrrolidinylmethyl)-2H-chromen-7-amine
obtained in Reference Example 107.

^1H NMR (CDCl_3) δ : 1.76 (4H, s), 2.01 (3H, s), 2.42 (4H, t, $J=5.1$ Hz), 2.49 (4H, s), 3.22 (2H, s), 3.48 (4H, t, $J=5.1$ Hz), 4.24 (1H, s), 4.68 (2H, s), 6.23 (1H, s), 6.77 (1H, s), 6.96 (1H, d, $J=8.7$ Hz), 7.09 (1H, d, $J=8.7$ Hz), 7.19-7.61 (9H, m).

Melting point: 104-106 °C (crystallization solvent: ethyl acetate - n-hexane)

Example 263

N-(2,2-Diphenylethyl)-N'-[4-methyl-3-(1-pyrrolidinylmethyl)-2H-chromen-7-yl]urea



The titled compound was obtained by carrying out the
same operation as in Reference Example 99, using 4-
methyl-3-(1-pyrrolidinylmethyl)-2H-chromen-7-amine

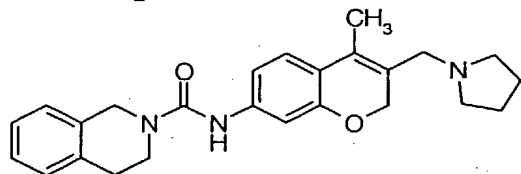
obtained in Reference Example 107.

¹H NMR (CDCl₃) δ: 1.76 (4H, s), 1.99 (3H, s), 2.49 (4H, s), 3.22 (2H, s), 3.83 (2H, t, J=7.8 Hz), 4.18 (1H, t, J=7.8 Hz), 4.66 (2H, s), 4.96 (1H, s), 6.48 (1H, s), 6.57 (1H, s), 6.69 (1H, d, J=8.1 Hz), 6.98 (1H, d, J=8.1 Hz), 7.20-7.30 (10H, m).

Melting point: 166-168 °C (crystallization solvent: ethyl acetate - n-hexane)

10 Example 264

N-[4-Methyl-3-(1-pyrrolidinylmethyl)-2H-chromen-7-yl]-3,4-dihydro-2(1H)-isoquinolinecarboxamide



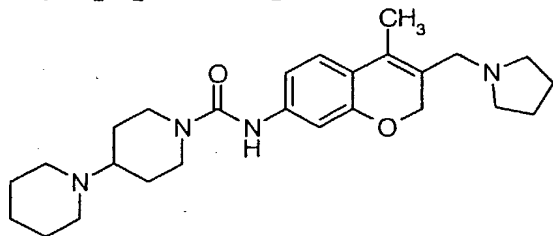
15 The titled compound was obtained by carrying out the same operation as in Reference Example 99, using 4-methyl-3-(1-pyrrolidinylmethyl)-2H-chromen-7-amine obtained in Reference Example 107.

¹H NMR (CDCl₃) δ: 1.76 (4H, s), 2.02 (3H, s), 2.49 (4H, s), 2.92 (2H, t, J=6.0 Hz), 3.23 (2H, s), 3.71 (2H, t, J=6.0 Hz), 4.65 (2H, s), 4.68 (2H, s), 6.43 (1H, s), 6.86 (1H, d, J=1.8 Hz), 7.02-7.22 (6H, m).

Melting point: 135-137 °C (crystallization solvent: ethyl acetate - n-hexane)

25 Example 265

N-[4-Methyl-3-(1-pyrrolidinylmethyl)-2H-chromen-7-yl]-4-(1-piperidiny)-1-piperidinecarboxamide

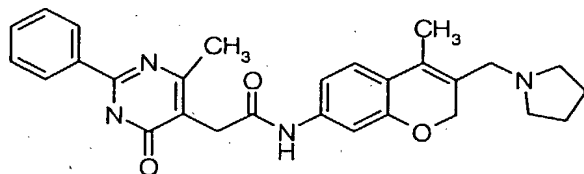


The titled compound was obtained by carrying out the same operation as in Reference Example 99, using 4-methyl-3-(1-pyrrolidinymethyl)-2H-chromen-7-amine obtained in Reference Example 107.

- 5 ^1H NMR (CDCl_3) δ : 1.27-1.89 (14H, m), 2.02 (3H, s), 2.49-2.51 (9H, m), 2.83-2.90 (2H, m), 3.23 (2H, s), 4.08-4.12 (2H, m), 4.68 (2H, s), 6.31 (1H, s), 6.80 (1H, d, $J=2.4$ Hz), 6.98 (1H, dd, $J=2.4$ Hz, 8.4 Hz), 7.09 (1H, d, $J=8.4$ Hz).
- 10 Melting point: 98-100 $^\circ\text{C}$ (crystallization solvent: ethyl acetate - n-hexane)

Example 266

- 2-(4-Methyl-6-oxo-2-phenyl-1,6-dihydro-5-pyrimidinyl)-
15 N-[4-methyl-3-(1-pyrrolidinymethyl)-2H-chromen-7-yl]acetamide

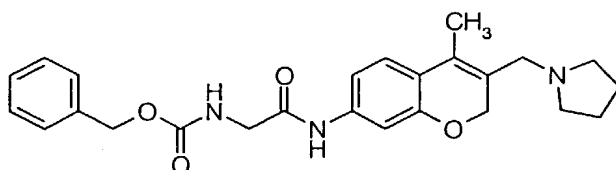


- The titled compound was obtained by carrying out the same operation as in Reference Example 1, using 4-methyl-3-(1-pyrrolidinymethyl)-2H-chromen-7-amine obtained in Reference Example 107.

- 20 ^1H NMR (CDCl_3) δ : 1.76 (4H, s), 1.98 (3H, s), 2.49 (4H, s), 2.61 (3H, s), 3.22 (2H, s), 3.65 (2H, s), 4.65 (2H, s), 6.86-7.00 (4H, m), 7.54 (3H, s), 8.01 (2H, s), 8.87 (1H, s).
- 25 Melting point: 255-257 $^\circ\text{C}$ (crystallization solvent: ethyl acetate - n-hexane)

Example 267

- Benzyl 2-[[4-methyl-3-(1-pyrrolidinymethyl)-2H-chromen-7-yl]amino]-2-oxoethylcarbamate
- 30



The titled compound was obtained by carrying out the same operation as in Reference Example 1, using 4-methyl-3-(1-pyrrolidinylmethyl)-2H-chromen-7-amine

5 obtained in Reference Example 107.

¹H NMR (CDCl₃) δ: 1.78 (4H, s), 2.03 (3H, s), 2.53 (4H, s), 3.26 (2H, s), 3.99 (2H, d, J=4.8 Hz), 4.71 (2H, s), 5.17 (2H, s), 5.50 (1H, bs), 7.00-7.14 (4H, m), 7.36 (5H, s), 7.80 (1H, bs).

10 Melting point: 143-145 °C (crystallization solvent: ethyl acetate - n-hexane)

Preparation Example 1

	(1) Compound obtained in	
15	Reference Example 25	50 mg
	(2) Lactose	34 mg
	(3) Corn starch	10.6 mg
	(4) Corn starch (paste)	5 mg
	(5) Magnesium stearate	0.4 mg
20	(6) Carboxymethylcellulose calcium	20 mg
	Total	120 mg

In accordance with a conventional manner, the above (1) to (6) are admixed and tableted using a tableting machine to give tablets.

25

Preparation Example 2

	(1) Compound obtained in Example 1	50 mg
	(2) Lactose	34 mg
	(3) Corn starch	10.6 mg
30	(4) Corn starch (paste)	5 mg
	(5) Magnesium stearate	0.4 mg
	(6) Carboxymethylcellulose calcium	20 mg
	Total	120 mg

In accordance with a conventional manner, the above (1) to (6) are admixed and tableted using a tableting machine to give tablets.

5 Reference Example 1-1

Amplification of rat SLC-1 receptor cDNA by PCR method using rat-brain-originated cDNA

Reverse transcription reaction was done using random primer, with rat-brain-originated poly (A)⁺RNA (Clone Tech Co.) used as a template. Reagent from the TaKaRa RNA PCR ver. 2 kit was used for the reverse transcription reaction.

Next, using this reverse transcription product as a template, amplification was done by a PCR method using synthetic DNA primers with sequence numbers 1 and 2.

15 Synthetic DNA primer was constructed to amplify genes in the domain where genes are translated by receptor protein.

At that time, individual restriction enzyme recognition sequences were also added on the 5' side and 3' side of the gene, to add a nucleotide sequence on the 5' side of gene which recognized restriction enzyme Sal I, and to add a nucleotide sequence on the 3' side of the gene which recognized the restriction enzyme Spe I. The reactant was constituted of 5 μ l of cDNA template, 0.4 μ M of synthetic DNA primer, 0.25 mM of dNTPs, 0.5 μ l of Pfu (StrataGene Co.) DNA polymerase, and buffers attached to enzymes, with total reaction quantity set at 50 μ l.

A thermal cycler (Parkin Elmer Co.) was used to produce cycles for amplification. After heating at 94°C for 60 seconds, the cycle consisting of 94°C for 60 seconds, 60°C for 30 seconds, and 72°C for 150 seconds, was repeated 35 times, and finally reaction was conducted at 72°C for 10 minutes. After 0.8% agarose gel electrophoresis, the amplified products were confirmed by ethidium bromide

35

Reference Example 1-2

Subcloning of PCR products into plasmid vector, and confirmation of an amplified cDNA sequence by decoding of a nucleotide sequence in an inserted cDNA portion

5 The reaction product after PCR conducted in Reference Example 1-1 was separated using 0.8% low-melting point agarose gel. After the band section was cut out using a razor, DNA was recovered by conducting fragmentation, phenol extraction, phenol-chloroform extraction and ethanol precipitation. The recovered DNA was subcloned on
10 plasmid vector PCR-Script Amp SK(+) in accordance with prescription of the PCR-Script™ Amp SK(+) cloning kit (Stratagene Co.). After this was introduced into Escherichia coli XL-1 Blue (Stratagene Co.) by transformation, the clones with fragments of inserted cDNA
15 were selected in LB agar culture medium containing ampicillin and X-gal. Only clones showing white color were separated using a sterilized toothpick, and transformant E. coli XL-1 Blue/rat SLC-1 was obtained.

Each clone was cultured overnight in LB culture medium
20 containing ampicillin, and plasmid DNA was prepared using QIA prep8 mini prep (Qiagen). A portion of the prepared DNA was digested with Sal I and Spe I, and the size of the inserted receptor cDNA fragment was confirmed. Reactions to determine nucleotide sequences were carried out using
25 a DyeDeoxy Terminator Cycle Sequence Kit (Parkin Elmer Co.), and decoded using a fluorescent light automatic sequencer. The sequences of the 3 clones obtained were analyzed, and it was confirmed that all of them match the reported gene sequence (Sequence number: 4) in which the
30 Sal I recognition sequence is added on the 5' side and the Spe I recognition sequence is added on the 3' side of the cDNA sequence (Lakaye, B., et al., Biochim. Biophys. Acta, Vol. 1401, pp. 216-220 (1998), accession No. AF08650) coding rat SLC-1 protein (Sequence number: 3).

35

Reference Example 1-3

Preparation of CHO cells for rat SLC-1 expression

The full-length amino acid sequence of rat brain originated SLC-1, which was confirmed in Reference Example 1-2, was coded, and plasmid was prepared using a plasmid
5 Midi Kit (Qiagen) from the E. coli transformed by the plasmid, to which the gene with Sal I recognition sequence added to the 5' side and Spe I recognition sequence added to the 3' side, had been introduced. Then, the insert
10 section was cut out by digesting with Sal I and Spe I. The insert DNA was cut out with a razor from the agarose gel after electrophoresis.

Next, fragmentation, phenol extraction, phenol-chloroform extraction, and ethanol precipitation, were conducted and the DNA was recovered. This insert DNA was
15 added to vector plasmid pAKKO-111H (the same vector plasmid as pAKKO1.11H described in Hinuma, S., et al., Biochim. Biophys. Acta, Vol. 1219, pp. 251-259 (1994)) for animal cell expression which was digested with Sal I and Spe I, and ligation was conducted using T4 ligase (TaKaRa Shuzo),
20 to construct pAKKO-SLC-1 plasmid for protein expression.

After E. coli DH5 transformed by pAKKO-SLC-1 was cultured, pAKKO-SLC-1 plasmid DNA was prepared using a Plasmid Midi Kit (Qiagen). This was introduced into CHO dhfr⁻ cells in accordance with the attached protocol, using
25 a CellPfect Transfection Kit (Amersham Pharmacia Biotech Co.). A coprecipitating suspension of 10 µg of DNA and calcium phosphate was prepared, and this suspension was added to 10 cm Petri dishes in which 5×10^5 or 1×10^6 of CHO dhfr⁻ cells had been seeded 24 hours previously. After
30 these cells were cultured for 1 day in MEMa culture medium containing 10% fetal bovine serum, subculture was conducted, and cultivation was conducted in selective culture medium, MEMa culture medium containing no nucleic acid but containing 10% dialyzed fetal bovine serum. 56
35 clones of colonies of the transformed CHO cells expressing SLC-1, proliferated in the selective culture medium, were

selected.

Reference Example 1-4

Selection of CHO/SLC-1 cell strain expressing a large
5 quantity of full-length rat SLC-1 receptor protein mRNA

The quantity of expressed full-length rat SLC-1
receptor protein mRNA of 56 clones of the CHO/SLC-1 strains
established in Reference Example 1-3, was measured using
a Cytostar T Plate (Amersham Pharmacia Biotech Co.) as shown
10 below according to the attached protocol. Each well of the
Cytostar T Plate was seeded with each clone of the CHO/SLC-1
strain by 2.5×10^4 , and cultured for 24 hours, then the
cells were fixed using 10% formalin. After 0.25% Triton
X-100 was added to each well to increase cell permeability,
15 ^{35}S -labeled riboprobes with sequence number: 5 were added
and hybridized. 20 mg/ml of RNaseA was added to each well
to digest free riboprobes. After the plate was thoroughly
washed, the radioactivity of the hybridized riboprobes was
determined using a Topcounter. Strains with high
20 radioactivity showed large amounts of mRNA expression. In
particular, mainly used was Clone number 44 among 3 clones
which showed large amounts of mRNA expression.

Reference Example 1-5

25 Isolation of plasmid containing human SLC-1 cDNA

After nicks were inserted into the DNA of Human fetal
brain originated cDNA library (SUPERSSCRIPT™ cDNA Library;
GIBCOBRL Co.) according to the manual of the Genetrapp
cDNA positive selection system (GIBCOBRL Co.), using phage
30 F1 endonuclease, single stranded human fetal brain
originated cDNA library was prepared by digesting the
above-mentioned library with Escherichia coli exonuclease
III.

Biotin-14-dCTP was added to the 3' end of synthetic
35 oligonucleotide (equivalent to 1434-1451 of accession No.
U71092), sequence number: 6 which was prepared according

to the report by Kolakowski Jr., et al. (Kolakowski Jr.,
et al. (1996) FEBS Lett. Vol. 398, pp. 253-258) using
Terminal Deoxynucleotidyl Transferase, and biotinated
oligonucleotide was prepared. The above manual was
5 followed regarding composition of a reaction mixture and
reaction time.

After 4 µg of single stranded human fetal brain
originated cDNA library was kept at 95°C for 1 minute, the
library was rapidly cooled on ice. 20 ng of biotinated
10 oligonucleotide was added, which was hybridized using the
attached hybridization buffer at 37°C for 1 hour.
Streptoavidin beads were added to the mixture, then single
stranded human fetal brain originated cDNA hybridized by
biotinated oligonucleotide, was isolated using a MAGNA-
15 SEP Magnetic Particle Separator (GIBCOBRL Co.). The
complementary strand was synthesized according to the
manual, using as primer 50 ng of synthetic oligonucleotide
(equivalent to 1011 - 1028 of accession No. U71092) of
sequence number: 7, prepared based on the report by
20 Kolakowski Jr., et al (Kolakowski Jr., et al. (1996) FEBS
Lett. Vol. 398, pp. 253-258), to give the double stranded
plasmid.

Reference Example 1-6

25 Determination of nucleotide sequence of plasmid containing
isolated human SLC-1 cDNA

After the plasmid obtained in Reference Example 1-
5 was introduced into ELECTROMAX™DH10B™ Cells by the
electroporation method, clones with cDNA inserted
30 fragments were selected in LB agar culture medium
containing ampicillin and X-gal. Using a sterilized
toothpick, only the clones showing white color were
separated to give transformant E. coli DH10B/hSLC-1.
Individual clones were cultured overnight in LB culture
35 medium containing ampicillin, and the plasmid DNA was
refined using QIA prep8 mini prep (Qiagen). The reactions

to determine nucleotide sequence were conducted using a DyeDeoxy Terminator Cycle Sequence Kit (Parkin Elmer Co.), and the nucleotide sequence was decoded using a fluorescent light automatic sequencer.

5 As the results, obtained was the sequence shown in Sequence number: 8. The amino acid sequence (Sequence number: 9) coded by the nucleotide sequence obtained here, differs from the human SLC-1 amino acid sequence predicted as the sequence analogized from rat SLC-1 based on human
10 chromosome DNA sequence (accession number: Z86090) containing human SLC-1 sequence, in the report by Lakaye, et al. (Lakaye, B., et al. (1998) *Biochim. Biophys. Acta*. Vol. 1401, pp. 216-220). This shows the presence of ATG, the initiation codon, on mRNA, in the 69 and 64 amino acids
15 upstream from the estimated sequence. *Escherichia coli* DH10B/phSLC1L8, the transformant produced by the plasmid containing DNA coding this sequence was deposited at IFO and NIBH.

20 Reference Example 1-7

Amplification of human SLC-1cDNA by PCR method using human fetal brain originated cDNA

Amplification by the PCR method was conducted using as the template plasmid containing human SLC-1 DNA sequence
25 cloned by the gene trap method, and using synthetic DNA primers of sequence number: 10 and sequence number: 11, and synthetic DNA primers of sequence number: 12 and sequence number: 13, respectively. The former amplified DNA and the latter amplified DNA were named as "human SLC-1(S)" and
30 "human SLC-1(L)", respectively. The synthetic DNA primer was constructed so that the genes in the domain translated to the receptor protein were amplified. At that time, a recognition sequence for each restriction enzyme was added on the 5' side and 3' side, so that the nucleotide sequence
35 recognized by restriction enzyme Sal I would be added on the 5' side of the gene, and the nucleotide sequence

recognized by restriction enzyme Spe I would be added on the 3' side. The composition of the reaction mixture for human SLC-1(S) amplification was: 5 µl of plasmid template containing human SLC-1 DNA sequence, 0.4 µM of respective synthetic DNA primers, 0.2 mM of dNTPs and 0.5 µl of Pfu DNA polymerase and buffers attached to the enzyme, with total quantity for reaction set at 50 µl. A thermal cycler (Parkin Elmer Co.) was used for the cycles for amplification. After heating at 94°C for 60 seconds, the cycle consisting of 94°C for 60 seconds, 57°C for 60 seconds, and 72°C for 150 seconds, was repeated 25 times, and finally the temperature of the reactant was maintained at 72°C for 10 minutes. The composition of the reaction mixture for human SLC-1(L) amplification was 5 µl of plasmid template containing human SLC-1 DNA sequence, 0.4 µM of respective synthetic DNA primers, 0.2 mM of dNTPs, 0.5 µl of Pfu DNA polymerase and buffers attached to the enzymes, with total quantity for reaction set at 50 µl. A thermal cycler (Parkin Elmer Co.) was used for the cycles for amplification. After heating at 94°C for 60 seconds, the cycle consisting of 94°C for 60 seconds, 60°C for 60 seconds, and 72°C for 3 minutes, was repeated 25 times, and finally the temperature of the reactant was maintained at 72°C for 10 minutes. After 0.8% agarose gel electrophoresis, confirmation of amplified products was conducted by ethidium bromide dying.

Reference Example 1-8

Subcloning of PCR product into plasmid vector and confirmation of amplified cDNA sequence by decoding of nucleotide sequence of inserted cDNA section

The reaction product after PCR in Reference Example 1-7 was separated using 0.8% low-melting point agarose gel, and the band section was cut out using a razor. After that, fragmentation, phenol extraction, phenol-chloroform extraction, and ethanol precipitation were conducted, and

the DNA was recovered. The recovered DNA was subcloned into pCR-Script Amp SK(+) plasmid vector, as prescribed by the PCR-Script™ Amp SK(+) cloning kit (Stratagene Co.). After this was introduced into Escherichia coli DH5a competent cells (TOYOBO) and transformed, the clones with cDNA inserted fragments were selected in LB agar culture medium containing ampicillin and X-gal. Using a sterilized toothpick, only clones showing white color were separated to give E. coli DH5α/hSLC-1(S), which is a transformant of human SLC-1 (S), and E. coli DH5α/hSLC-1(L), which is a transformant of human SLC-1 (L). Each clone was cultured overnight in LB culture medium containing ampicillin, and plasmid DNA was prepared using QIA prep8 mini prep (Qiagen).

Some of the prepared DNA was digested with Sal I and Spe I restriction enzymes, and the size of the receptor cDNA fragments inserted was confirmed. The reactions to determine nucleotide sequence were conducted using a DyeDeoxy Terminator Cycle Sequence Kit (Parkin Elmer Co.) and the nucleotide sequence was decoded using a fluorescent light automatic sequencer. The sequence of the obtained clones respectively matched the DNA sequence (sequence number:14) which should be amplified by synthetic DNA primers of sequence number: 10 and sequence number: 11 using human SLC-1 gene as a template, and the DNA sequence (sequence number: 15) which should be amplified by synthetic DNA primers of sequence number: 12 and sequence number: 13 using human SLC-1 gene as a template.

Reference Example 1-9

Preparation of CHO cells for expression of human SLC-1(S), and CHO cells for expression of human SLC-1(L)

Plasmid was prepared from the E. coli clones transformed by the plasmid wherein inserted were human SLC-1(S) and human SLC-1(L) whose sequences were confirmed in Reference Example 1-8, using a Plasmid Midi Kit (Qiagen), and the insert section was cut out using Sal I and Spe I

restriction enzymes. After electrophoresis was conducted, the insert DNA was cut out from agarose gel using a razor. Next, fragmentation, phenol extraction, phenol-chloroform extraction, and ethanol precipitation were conducted, and the insert DNA was recovered.

This insert DNA was added to pAKKO-111H vector plasmid for animal cell expression, digested with Sal I and Spe I (the same vector plasmid as the pAKKO1.11H described in Hinuma, S., et al., Biochim. Biophys. Acta, Vol. 1219, pp. 251-259 (1994)), and ligation was conducted by adding T4 ligase (TaKaRa Shuzo), to construct pAKKO-hSLC-1(S) and pAKKO-hSLC-1(L) plasmids for protein expression.

After E. coli DH5 α (TOYOBO) transformed by pAKKO-hSLC-1(S) and pAKKO-hSLC-1(L) was cultured, pAKKO-hSLC-1(S) and pAKKO-hSLC-1(L) plasmid DNAs were prepared using a Plasmid Midi Kit (Qiagen). These were introduced into CHO dhfr⁻ cells in accordance with the attached protocol, using a CellPfect Transfection Kit (Amersham Pharmacia Biotech Co.). A coprecipitative suspension of 10 μ g of DNA with calcium phosphate was made, which was added to 10 cm Petri dishes seeded 24 hours in advance with 5×10^5 or 1×10^6 CHO dhfr⁻ cells. After the above was cultured for 1 day in MEM α culture medium containing 10% fetal bovine serum, subculture was conducted, and then cultivation was conducted in MEM α culture medium containing no nucleic acid but containing 10% dialyzed fetal bovine serum, which is a selective culture medium. 56 clones of colonies of transformed cells which are human SLC-1(S) gene introduced CHO cells, and 61 clones of colonies of transformed cells which are human SLC-1(L) gene introduced CHO cells, both of which proliferated in the selective culture medium, were selected.

Reference Example 1-10

Selection of cell colonies into which genes with large quantities of human SLC-1(S) and human SLC-1 (L) mRNA

expression have been introduced

The quantities of expressed mRNA of 56 clones of CHO/hSLC-1(S) colonies and 61 clones of CHO/hSLC-1(L) colonies, both of which were established in Reference Example 1-9, were measured in accordance with the attached
5 protocol using a Cytostar T Plate (Amersham Pharmacia Biotech Co.) as shown below.

After each well of the Cytostar T Plate was seeded with each clone of CHO/hSLC-1(S) colonies and CHO/hSLC-1(L)
10 colonies by 2.5×10^4 , and cultured for 24 hours, the cells were fixed using 10% formalin.

After 0.25% Triton X-100 was added to each well to increase cell permeability, ^{35}S -labeled riboprobe of sequence number: 16 was added and hybridization was
15 conducted.

20 mg/ml of RNaseA was added to each well to digest free riboprobe. After the plate was washed well, the radioactivity of the hybridized riboprobe was determined. Colonies showing high radioactivity expressed large
20 quantities of mRNA. Of the 7 clones which expressed large quantities of mRNA, mainly used was Clone number 57.

Experimental Example 1

Determination of antagonist activity using GTPgS binding
25 assay of test compound

Membrane fraction was prepared by the following method, using the human SLC-1 expressing CHO cell clone 57 obtained in Reference Example 1-10, and the rat SLC-1 expressing CHO cell clone 44 obtained in Reference Example
30 1-4.

The human and rat SLC-1 expressing CHO cells (1×10^8) were scraped in buffer saline phosphate (pH 7.4) to which 5 mM EDTA (ethylenediaminetetraacetic acid) had been added, and centrifuged. 10 ml of homogenized buffer (10 mM NaHCO_3 ,
35 5 mM EDTA, pH 7.5) was added to the cell pellets, and they were homogenized using a Polytron homogenizer. The

supernatant obtained by centrifugation at $400 \times g$ for 15 minutes was further centrifuged at $100,000 \times g$ for 1 hour, to obtain the membrane fraction precipitate. This precipitate was suspended in 2 ml of assay buffer [50 mM Tris-HCl(pH 7.5), 1 mM EDTA, 0.1% BSA (bovine serum albumin), 10 mM $MgCl_2$, 100 mM NaCl, 1 μ M GDP (guanosine 5'-diphosphate), 0.25 mM PMSF (phenylmethylsulfonyl fluoride), 1 mg/ml pepstatin, 20 mg/ml leupeptin, 10 mg/ml phosphoramidon], which was centrifuged at $100,000 \times g$ for 1 hour. The membrane fraction recovered as precipitate was suspended again in 2 ml of assay buffer, and after the suspension was divided, individual portions were preserved at $-80^\circ C$ and thawed before every use.

Determination of antagonist activity of the test compound was conducted as shown below. After 171 μ l of SLC-1 expressing CHO cell membrane fractions diluted with assay buffer was poured into each well of a 96-well polypropylene plate, 2 μ l of 3×10^{-10} M MCH diluted with DMSO solution, 2 μ l of test compound solution diluted to various concentrations, and 25 μ l of [^{35}S]-Guanosine 5'-(γ -thio) triphosphate (produced by Daiichi Kagaku Yakuhin) were added respectively. (Final concentration of cell membrane: 20 μ g/ml, final concentration of [^{35}S]-Guanosine 5'-(γ -thio) triphosphate: 0.33 nM).

After this reaction mixture was allowed to react for 1 hour under stirring, it was filtered under vacuum using a glass filter (GF-C), then the filter was washed 3 times with 300 μ l of washing solution (50 mM Tris-HCl buffer solution pH 7.5). 50 ml of liquid scintillator was added to the glass filter, and residual radioactivity was determined using a liquid scintillation counter.

The IC_{50} value of the compound was calculated from the binding inhibition rate (%), based on the definition that the binding inhibition rate (%) = (radioactivity when compound and MCH were added - radioactivity when DMSO solution was added)/(radioactivity when MCH was added -

radioactivity when DMSO solution was added) $\times 100$.

The results were shown below.

Compound Number	Inhibition Activity (IC ₅₀ value: nM)
Reference Example 25	90
Example 1	40

5

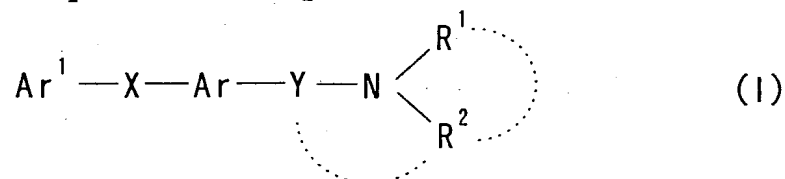
Industrial Applicability

Compounds (I), (I') and salts thereof possess excellent MCH receptor antagonistic activities, and are useful as an agent for preventing or treating obesity, etc.

10

CLAIMS

1. A melanin-concentrating hormone antagonist which comprises a compound of the formula :



5 wherein Ar¹ is a cyclic group which may have substituents; X is a spacer having a main chain of 1 to 6 atoms; Y is a bond or a spacer having a main chain of 1 to 6 atoms; Ar is a monocyclic aromatic ring which may be condensed with
10 a 4 to 8 membered non-aromatic ring, and may have further substituents;

R¹ and R² are independently hydrogen atom or a hydrocarbon group which may have substituents; R¹ and R², together with the adjacent nitrogen atom, may form a nitrogen-containing
15 hetero ring which may have substituents; R² may form a spiro ring together with Ar; or R², together with the adjacent nitrogen atom and Y, may form a nitrogen-containing hetero ring which may have substituents; or a salt thereof.

20 2. An antagonist according to claim 1, wherein Y is a spacer having a main chain of 1 to 6 atoms; R¹ and R² are independently hydrogen atom or a hydrocarbon group which may have substituents; R¹ and R², together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring
25 which may have substituents; or R² may form a spiro ring together with Ar.

3. An antagonist according to claim 2, wherein Ar¹ is an aromatic group which may have substituents; and "a hydrocarbon group which may have substituents" for R¹ and
30 R² is "C₁₋₆ alkyl which may have substituents".

4. An antagonist according to claim 1, wherein the cyclic

group for Ar¹ is C₆₋₁₄ monocyclic or condensed polycyclic aromatic hydrocarbon group.

5. An antagonist according to claim 1, wherein the cyclic group for Ar¹ is a group formed by removing an optional one hydrogen atom from an aromatic ring assemble in which 2 or 3 C₆₋₁₄ monocyclic or condensed polycyclic aromatic hydrocarbon groups are directly bonded by single bonds.

6. An antagonist according to claim 1, wherein the cyclic group for Ar¹ is a group formed by removing an optional one hydrogen atom from an aromatic ring assemble in which C₆₋₁₄ monocyclic or condensed polycyclic aromatic hydrocarbon and 5 to 10 membered aromatic hetero ring are directly bonded by a single bond.

7. An antagonist according to claim 1, wherein Ar¹ is phenyl, biphenyl, phenyl-pyridyl, phenyl-furyl, phenyl-isoxazolyl, diphenyl-oxazolyl, pyridyl-phenyl, phenyl-pyrimidinyl, benzofuranyl-phenyl, furyl-phenyl, terphenyl, thienyl-phenyl, indolyl, naphthyl-oxadiazolyl, benzofuranyl-oxadiazolyl, benzothienyl, benzofuranyl, fluorenyl, pyridyl-pyrrolyl or thioxanthanyl;
each of which may have 1 to 3 substituents selected from the group consisting of halogen atom; nitro; C₁₋₃ alkylenedioxy; optionally halogenated C₁₋₆ alkyl; hydroxy-C₁₋₆ alkyl; optionally halogenated C₃₋₆ cycloalkyl; optionally halogenated C₁₋₆ alkoxy; optionally halogenated C₁₋₆ alkythio; hydroxy; C₇₋₁₉ aralkyloxy which may have substituents; C₆₋₁₄ aryloxy which may have substituents; amino; mono-C₁₋₆ alkylamino; di-C₁₋₆ alkylamino; 5 to 7 membered saturated cyclic amino which may have substituents and may be condensed with a benzene ring; 5 to 7 membered non-aromatic heterocyclic groups which may have substituents; formyl; carboxy; C₆₋₁₄ aryl-carbonyl which may

have substituents; C₆₋₁₄ aryl-carbamoyl which may have substituents; aromatic hetero ring-carbamoyl which may have substituents; C₁₋₆ alkoxy-carbonyl; optionally halogenated C₁₋₆ alkyl-carboxamide; C₆₋₁₄ aryl-carboxamide which may have substituents; C₇₋₁₉ aralkyl-carboxamide which may have substituents; aromatic hetero ring-carboxamide which may have substituents; N-(C₆₋₁₄ aryl-carbonyl which may have substituents)-N-C₁₋₆ alkylamino; C₆₋₁₄ arylamino-carbonylamino which may have substituents; C₆₋₁₄ arylsulfonylamino which may have substituents; C₆₋₁₄ aryl-carbonyloxy which may have substituents; oxo; carboxy-C₁₋₆ alkyl; C₁₋₆ alkoxy-carbonyl-C₁₋₆ alkyl; C₇₋₁₉ aralkyl which may have substituents; aromatic hetero ring-C₁₋₆ alkoxy; and cyano.

8. An antagonist according to claim 1, wherein Ar¹ is piperidinyl, piperazinyl, pyrrolidinyl, dihydropyridyl or tetrahydropyridyl; each of which may have 1 or 2 substituents selected from the group consisting of oxo, C₆₋₁₄ aryl which may have substituents, hydroxy, C₇₋₁₉ aralkyloxy-carbonyl, and C₇₋₁₉ aralkyl.

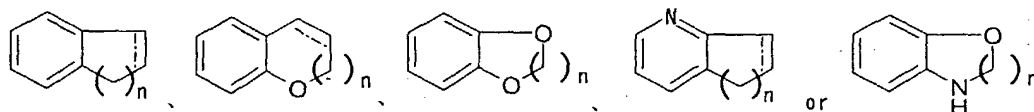
9. An antagonist according to claim 1, wherein the "spacer having a main chain of 1 to 6 atoms" for X and Y is a bivalent group consisting of 1 to 3 species selected from -O-, -S-, -CO-, -SO-, -SO₂-, -NR⁸- (R⁸ is hydrogen atom, optionally halogenated C₁₋₆ alkyl, optionally halogenated C₁₋₆ alkyl-carbonyl, optionally halogenated C₁₋₆ alkylsulfonyl), and a bivalent C₁₋₆ non-cyclic hydrocarbon group which may have substituents.

10. An antagonist according to claim 1, wherein X is -CONR^{8c}-, -NR^{8c}CO-, -CH=CH-CONR^{8c}- or -SO₂NR^{8c}- wherein R^{8c} is hydrogen atom or C₁₋₆ alkyl.

11. An antagonist according to claim 1, wherein Y is an

optionally halogenated bivalent C_{1-6} non-cyclic hydrocarbon group.

12. An antagonist according to claim 1, wherein Ar is a
5 ring of the formula :



wherein ----- is a single bond or double bond, n is an integer of 1 to 4.

- 10 13. An antagonist according to claim 1, wherein R^1 and R^2 are hydrogen atom or C_{1-6} alkyl which may have substituents; or R^1 and R^2 , together with the adjacent nitrogen atom, form a 3 to 8 membered nitrogen-containing hetero ring.

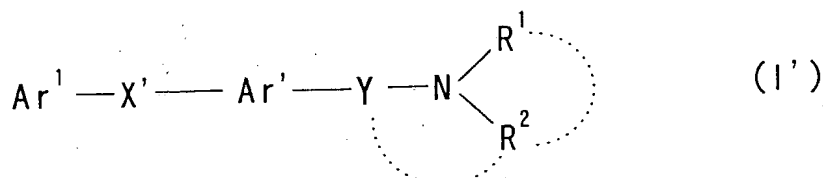
- 15 14. An antagonist according to claim 1, which is an agent for preventing or treating diseases caused by a melanin-concentrating hormone.

- 20 15. An antagonist according to claim 1, which is an agent for preventing or treating obesity.

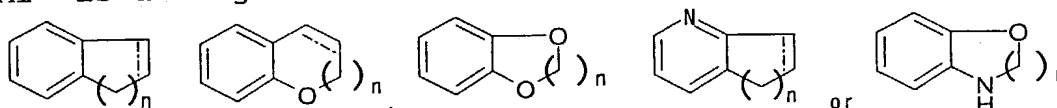
16. An antagonist according to claim 1, which is an anorectic agent.

- 25 17. A pharmaceutical, which comprises a melanin-concentrating hormone antagonist in combination with at least one species selected from the group consisting of an agent for treating diabetes, an agent for treating hypertension and an agent for treating arteriosclerosis.

- 30 18. A compound of the formula :

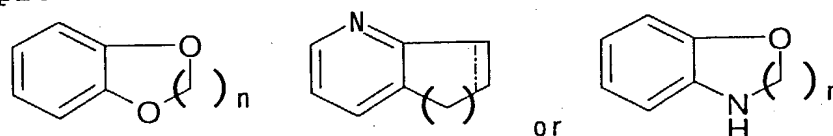


wherein Ar^1 is a cyclic group which may have substituents;
 Ar' is a ring of the formula :



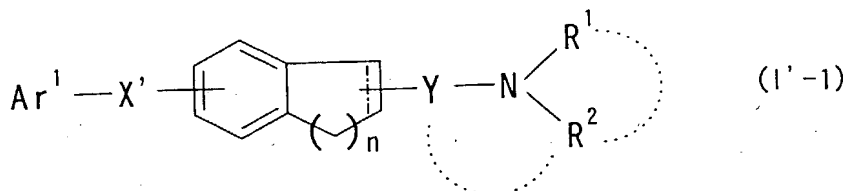
- 5 wherein ----- is a single bond or double bond, n is an integer of 1 to 4, and each ring may have substituents;
 X' is $-\text{CONR}^{\text{sc}}-$, $-\text{NR}^{\text{sc}}\text{CO}-$, $-\text{CH}=\text{CH}-\text{CONR}^{\text{sc}}-$ or $-\text{SO}_2\text{NR}^{\text{sc}}-$ where R^{sc} is hydrogen atom or C_{1-6} alkyl;
 Y is a spacer having a main chain of 1 to 6 atoms;
 10 R^1 and R^2 are independently hydrogen atom or a hydrocarbon group which may have substituents; R^1 and R^2 , together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; or R^2 , together with the adjacent nitrogen atom and Y , may form a
 15 nitrogen-containing hetero ring which may have substituents;

provided that Ar' is a ring of the formula :



- wherein symbols have the same meanings as defined above,
 20 and each ring may have substituents, when X' is $-\text{SO}_2\text{NH}-$; and provided that Ar^1 is not biphenyl which may be substituted, when X' is $-\text{CONH}-$ and Ar' is any one of benzopyran, dihydrobenzopyran, dihydrobenzoxazine, dihydrobenzoxazole or tetrahydrobenzoxazepine;
 25 (excluding $\text{N}-[2-(\text{N},\text{N}-\text{dimethylamino})\text{methyl}-6\text{-tetralinyl}]-4\text{-biphenylcarboxamide}$); or a salt thereof.

19. A compound of the formula :



wherein Ar¹ is a cyclic group which may have substituents;

----- is a single bond or double bond;

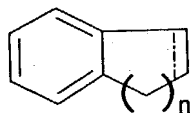
n is an integer of 1 to 4;

5 X' is -CONR^{8c}-, -NR^{8c}CO- or -CH=CH-CONR^{8c}- where R^{8c} is hydrogen atom or C₁₋₆ alkyl;

Y is a spacer having a main chain of 1 to 6 atoms;

R¹ and R² are independently hydrogen atom or a hydrocarbon group which may have substituents; R¹ and R², together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; or R², together with the adjacent nitrogen atom and Y, may form a nitrogen-containing hetero ring which may have substituents;

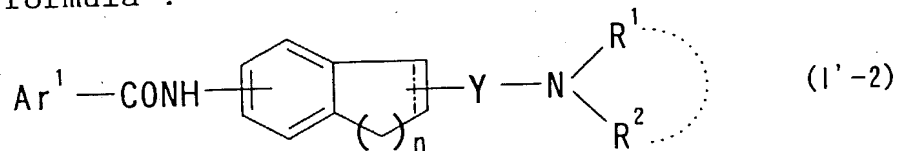
15 a ring of the formula :



wherein symbols have the same meanings as defined above, may have further substituents;

provided that N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]-4-biphenylcarboxamide is excluded; or a salt thereof.

20. A compound according to claim 19, which is of the formula :



25

wherein R¹ and R² are independently hydrogen atom or a hydrocarbon group which may have substituents; R¹ and R²,

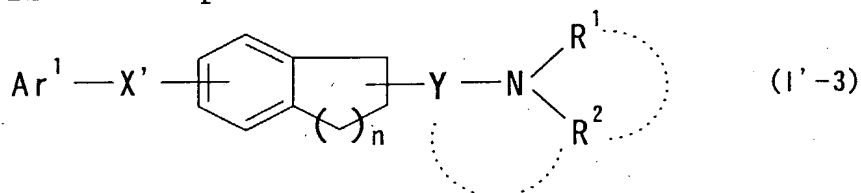
together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; the other symbols have the same meanings as defined in claim 19.

5

21. A compound according to claim 20, wherein Ar^1 is an aromatic group which may have substituents; and "a hydrocarbon group which may have substituents" for R^1 and R^2 is " C_{1-6} alkyl which may have substituents".

10

22. A compound of the formula :



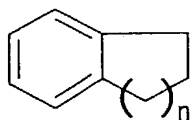
wherein Ar^1 is a cyclic group which may have substituents; n is an integer of 1 to 4;

15 X' is $-\text{CONR}^{\text{sc}}-$, $-\text{NR}^{\text{sc}}\text{CO}-$ or $-\text{CH}=\text{CH}-\text{CONR}^{\text{sc}}-$ where R^{sc} is hydrogen atom or C_{1-6} alkyl;

Y is a spacer having a main chain of 1 to 6 atoms;

R^1 and R^2 are independently hydrogen atom or a hydrocarbon group which may have substituents; R^1 and R^2 , together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; or R^2 , together with the adjacent nitrogen atom and Y , may form a nitrogen-containing hetero ring which may have substituents;

25 a ring of the formula :



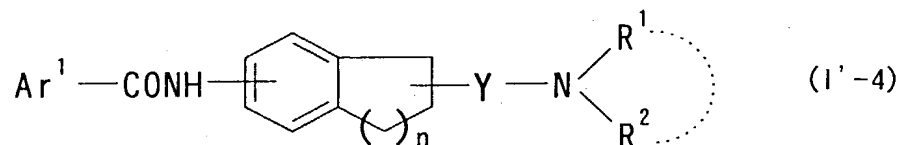
wherein n has the same meaning as defined above, may have further substituents;

provided that N-[2-(N,N-dimethylamino)methyl-6-

30 tetralinyl]-4-biphenylcarboxamide is excluded; or a salt

thereof.

23. A compound according to claim 22, which is of the formula :



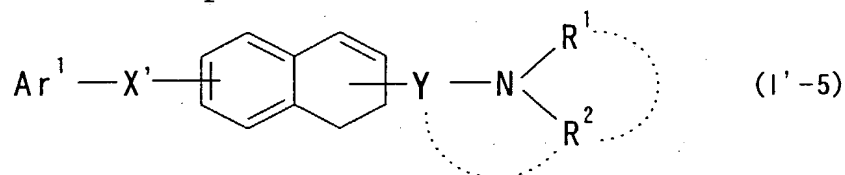
wherein R^1 and R^2 are independently hydrogen atom or a hydrocarbon group which may have substituents; R^1 and R^2 , together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; the other symbols have the same meanings as defined in claim 22.

10

24. A compound according to claim 23, wherein Ar^1 is an aromatic group which may have substituents; and "a hydrocarbon group which may have substituents" for R^1 and R^2 is " C_{1-6} alkyl which may have substituents".

15

25. A compound of the formula :

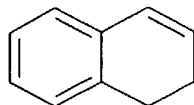


wherein Ar^1 is a cyclic group which may have substituents; X' is $-\text{CONR}^{\text{sc}}-$, $-\text{NR}^{\text{sc}}\text{CO}-$ or $-\text{CH}=\text{CH}-\text{CONR}^{\text{sc}}-$ where R^{sc} is hydrogen atom or C_{1-6} alkyl; Y is a spacer having a main chain of 1 to 6 atoms; R^1 and R^2 are independently hydrogen atom or a hydrocarbon group which may have substituents; R^1 and R^2 , together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; or R^2 , together with the adjacent nitrogen atom and Y , may form a nitrogen-containing hetero ring which may have substituents;

25

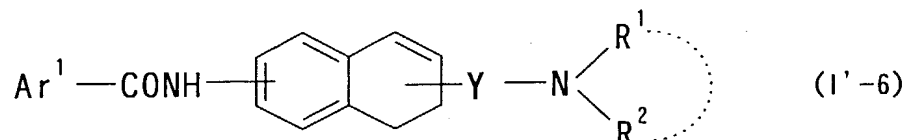
30

a ring of the formula :



may have further substituents; or a salt thereof.

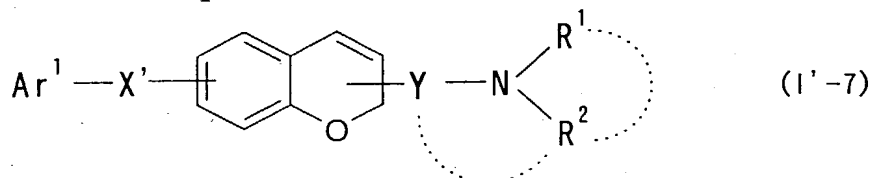
- 5 26. A compound according to claim 25, which is of the formula :



- 10 wherein R¹ and R² are independently hydrogen atom or a hydrocarbon group which may have substituents; R¹ and R², together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; the other symbols have the same meanings as defined in claim 25.

- 15 27. A compound according to claim 26, wherein Ar¹ is an aromatic group which may have substituents; and "a hydrocarbon group which may have substituents" for R¹ and R² is "C₁-₆ alkyl which may have substituents".

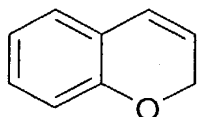
- 20 28. A compound of the formula :



- wherein Ar¹ is a cyclic group which may have substituents; X' is -CONR^{8c}-, -NR^{8c}CO-, -CH=CH-CONR^{8c}- or -SO₂NR^{8c}- where R^{8c} is hydrogen atom or C₁-₆ alkyl;
- 25 Y is a spacer having a main chain of 1 to 6 atoms; R¹ and R² are independently hydrogen atom or a hydrocarbon group which may have substituents; R¹ and R², together with the adjacent nitrogen atom, may form a nitrogen-containing

hetero ring which may have substituents; or R^2 , together with the adjacent nitrogen atom and Y, may form a nitrogen-containing hetero ring which may have substituents;

5 a ring of the formula :

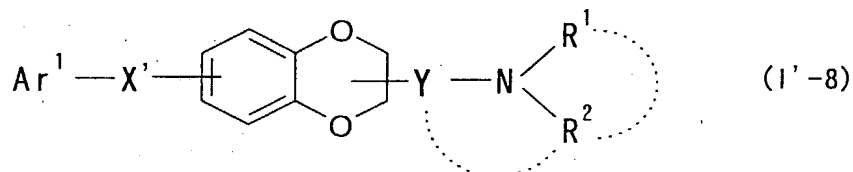


may have further substituents;

provided that Ar^1 is not biphenyl which may be substituted, when X' is $-CONH-$; or a salt thereof.

10

29. A compound of the formula :



wherein Ar^1 is a cyclic group which may have substituents; X' is $-CONR^{8c}-$, $-NR^{8c}CO-$, $-CH=CH-CONR^{8c}-$ or $-SO_2NR^{8c}-$ where R^{8c} is hydrogen atom or C_{1-6} alkyl;

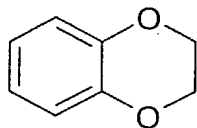
15

Y is a spacer having a main chain of 1 to 6 atoms;

R^1 and R^2 are independently hydrogen atom or a hydrocarbon group which may have substituents; R^1 and R^2 , together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; or R^2 , together with the adjacent nitrogen atom and Y, may form a nitrogen-containing hetero ring which may have substituents;

20

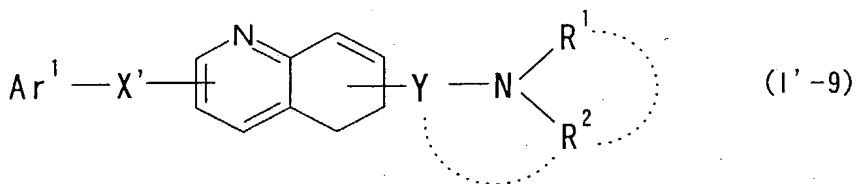
a ring of the formula :



25

may have further substituents; or a salt thereof.

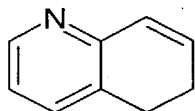
30. A compound of the formula :



wherein Ar¹ is a cyclic group which may have substituents;
 X' is -CONR^{8c}-, -NR^{8c}CO-, -CH=CH-CONR^{8c}- or -SO₂NR^{8c}- where
 R^{8c} is hydrogen atom or C₁₋₆ alkyl;

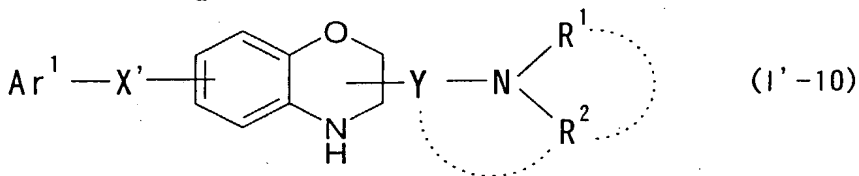
- 5 Y is a spacer having a main chain of 1 to 6 atoms;
 R¹ and R² are independently hydrogen atom or a hydrocarbon
 group which may have substituents; R¹ and R², together with
 the adjacent nitrogen atom, may form a nitrogen-containing
 hetero ring which may have substituents; or R², together
 10 with the adjacent nitrogen atom and Y, may form a
 nitrogen-containing hetero ring which may have
 substituents;

a ring of the formula :



- 15 may have further substituents; or a salt thereof.

31. A compound of the formula :

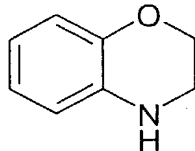


wherein Ar¹ is a cyclic group which may have substituents;
 20 X' is -CONR^{8c}-, -NR^{8c}CO-, -CH=CH-CONR^{8c}- or -SO₂NR^{8c}- where
 R^{8c} is hydrogen atom or C₁₋₆ alkyl;

- Y is a spacer having a main chain of 1 to 6 atoms;
 R¹ and R² are independently hydrogen atom or a hydrocarbon
 group which may have substituents; R¹ and R², together with
 25 the adjacent nitrogen atom, may form a nitrogen-containing
 hetero ring which may have substituents; or R², together
 with the adjacent nitrogen atom and Y, may form a
 nitrogen-containing hetero ring which may have

substituents;

a ring of the formula :



may have further substituents;

- 5 provided that Ar¹ is not biphenyl which may be substituted, when X' is -CONH-; or a salt thereof.

32. A pharmaceutical composition which comprises a compound as defined in any one of claims 18, 19, 22, 25,
10 26, 28, 29, 30 and 31.

33. A prodrug of a compound as defined in any one of claims 18, 19, 22, 25, 26, 28, 29, 30 and 31.

- 15 34. A compound according to claim 18, which is
N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]-(4'-methoxybiphenyl-4-yl)carboxamide;
4'-fluoro-N-[6-[(N,N-dimethylamino)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;
20 4'-fluoro-N-[6-(1-piperidinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;
4'-fluoro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;
25 (+)-4'-fluoro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;
(-)-4'-fluoro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;
30 4'-chloro-N-[3-[(N,N-dimethylamino)methyl]-2H-chromen-7-yl][1,1'-biphenyl]-4-carboxamide;
4'-fluoro-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-

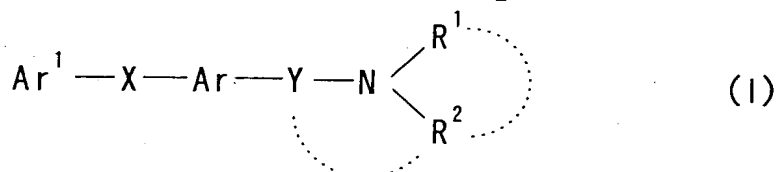
- naphthalenyl][1,1'-biphenyl]-4-carboxamide;
N-[3-[(dimethylamino)methyl]-2H-chromen-7-yl]-4'-fluoro[1,1'-biphenyl]-4-carboxamide;
4'-chloro-N-[6-[(dimethylamino)methyl]-5-methyl-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;
5 6-(4-methoxyphenyl)-N-[5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]nicotinamide;
4'-chloro-N-[7-[(dimethylamino)methyl]-5,6-dihydro-3-quinolinyl][1,1'-biphenyl]-4-carboxamide;
10 4-(4-chlorophenyl)-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]-3,6-dihydro-1(2H)-pyridinecarboxamide;
N-[6-[(dimethylamino)methyl]-7,8-dihydro-2-naphthalenyl]-4-(4-fluorophenyl)-1-piperidinecarboxamide;
15 4-(4-methoxyphenyl)-N-[6-(1-pyrrolidinylmethyl)-5-methyl-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide;
20 4'-fluoro-N-[6-[2-(1-pyrrolidinyl)ethyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;
4'-chloro-N-[6-[2-(1-pyrrolidinyl)ethyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;
4'-chloro-N-[2-[(dimethylamino)methyl]-3,4-dihydro-2H-1,4-benzoxazin-6-yl][1,1'-biphenyl]-4-carboxamide;
25 4-(4-methoxyphenyl)-N-[5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide;
4-(4-chlorophenyl)-N-[6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide;
30 4'-chloro-N-[2-[(dimethylamino)methyl]-1H-inden-6-yl][1,1'-biphenyl]-4-carboxamide;
4'-fluoro-N-[2-(1-pyrrolidinylmethyl)-3,4-dihydro-2H-1,4-benzoxazin-6-yl][1,1'-biphenyl]-4-carboxamide;
35 4'-fluoro-N-[5-methyl-6-[(4-methyl-1-

piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;

4'-chloro-N-[5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide; or

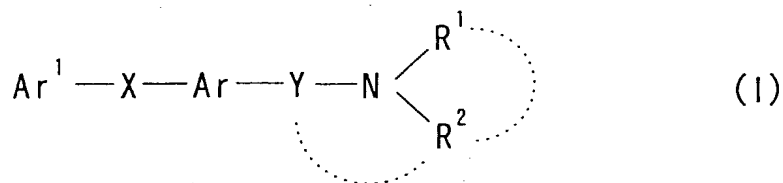
4-(4-chlorophenyl)-N-[5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide.

35. A method for preventing or treating diseases caused by a melanin-concentrating hormone in a mammal in need thereof, which comprises administering to said mammal an effective amount of a compound of the formula :



- wherein Ar^1 is a cyclic group which may have substituents; X is a spacer having a main chain of 1 to 6 atoms; Y is a bond or a spacer having a main chain of 1 to 6 atoms; Ar is a monocyclic aromatic ring which may be condensed with a 4 to 8 membered non-aromatic ring, and may have further substituents; R^1 and R^2 are independently hydrogen atom or a hydrocarbon group which may have substituents; R^1 and R^2 , together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; R^2 may form a spiro ring together with Ar; or R^2 , together with the adjacent nitrogen atom and Y, may form a nitrogen-containing hetero ring which may have substituents; or a salt thereof.

36. A method for preventing or treating obesity in a mammal in need thereof, which comprises administering to said mammal an effective amount of a compound of the formula :



wherein Ar^1 is a cyclic group which may have substituents;

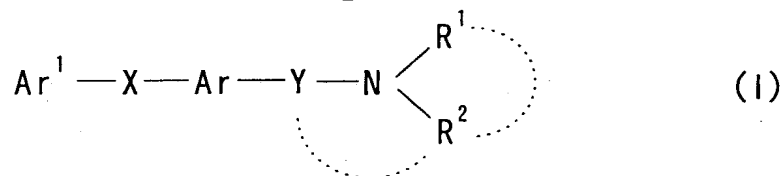
X is a spacer having a main chain of 1 to 6 atoms;

Y is a bond or a spacer having a main chain of 1 to 6 atoms;

Ar is a monocyclic aromatic ring which may be condensed with a 4 to 8 membered non-aromatic ring, and may have further substituents;

R^1 and R^2 are independently hydrogen atom or a hydrocarbon group which may have substituents; R^1 and R^2 , together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; R^2 may form a spiro ring together with Ar; or R^2 , together with the adjacent nitrogen atom and Y, may form a nitrogen-containing hetero ring which may have substituents; or a salt thereof.

37. Use of a compound of the formula :



wherein Ar^1 is a cyclic group which may have substituents;

X is a spacer having a main chain of 1 to 6 atoms;

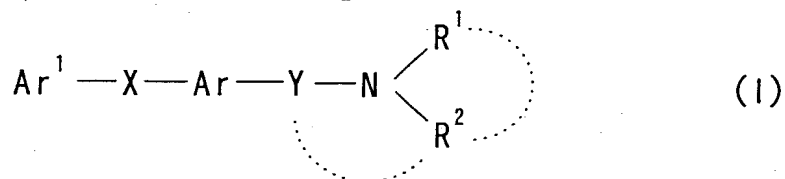
Y is a bond or a spacer having a main chain of 1 to 6 atoms;

Ar is a monocyclic aromatic ring which may be condensed with a 4 to 8 membered non-aromatic ring, and may have further substituents;

R^1 and R^2 are independently hydrogen atom or a hydrocarbon group which may have substituents; R^1 and R^2 , together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; R^2 may form a spiro ring together with Ar; or R^2 , together with the adjacent nitrogen atom and Y, may form a nitrogen-containing hetero ring which may have substituents; or a salt thereof;

for the manufacture of a pharmaceutical preparation for preventing or treating diseases caused by a melanin-concentrating hormone.

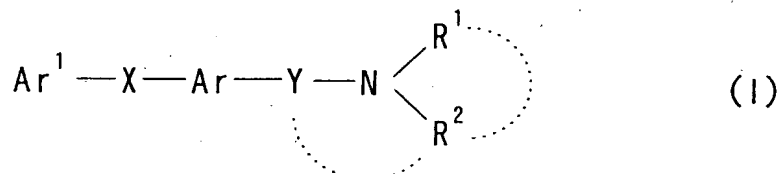
- 5 38. Use of a compound of the formula :



- wherein Ar^1 is a cyclic group which may have substituents;
X is a spacer having a main chain of 1 to 6 atoms;
Y is a bond or a spacer having a main chain of 1 to 6 atoms;
10 Ar is a monocyclic aromatic ring which may be condensed with a 4 to 8 membered non-aromatic ring, and may have further substituents;
 R^1 and R^2 are independently hydrogen atom or a hydrocarbon group which may have substituents; R^1 and R^2 , together with the adjacent nitrogen atom, may form a nitrogen-containing
15 hetero ring which may have substituents; R^2 may form a spiro ring together with Ar; or R^2 , together with the adjacent nitrogen atom and Y, may form a nitrogen-containing hetero ring which may have substituents; or a salt thereof;
20 for the manufacture of a pharmaceutical preparation for preventing or treating obesity.

ABSTRACT

5 A melanin-concentrating hormone antagonist which
comprises a compound of the formula :



wherein Ar¹ is a cyclic group which may have substituents;
X is a spacer having a main chain of 1 to 6 atoms;
Y is a bond or a spacer having a main chain of 1 to 6 atoms;
10 Ar is a monocyclic aromatic ring which may be condensed with
a 4 to 8 membered non-aromatic ring, and may have further
substituents;
R¹ and R² are independently hydrogen atom or a hydrocarbon
group which may have substituents; R¹ and R², together with
15 the adjacent nitrogen atom, may form a nitrogen-containing
hetero ring which may have substituents; R² may form a spiro
ring together with Ar; or R², together with the adjacent
nitrogen atom and Y, may form a nitrogen-containing hetero
ring which may have substituents; or a salt thereof;
20 which is useful as an agent for preventing or treating
obesity, etc.

Declaration and Power of Attorney for Patent Application

特許出願宣言書及び委任状

Japanese Language Declaration

日本語宣言書

私は、以下に記名された発明者として、ここに下記の通り宣言する：

As a below named inventor, I hereby declare that:

私の住所、郵便の宛先そして国籍は、私の氏名の後に記載された通りである。

My residence, post office address and citizenship are as stated next to my name.

下記の名称の発明について、特許請求範囲に記載され、且つ特許が求められている発明主題に関して、私は、最初、最先且つ唯一の発明者である（唯一の氏名が記載されている場合）か、或いは最初、最先且つ共同発明者である（複数の氏名が記載されている場合）と信じている。

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled

MELANIN CONCENTRATING

HORMONE ANTAGONIST

上記発明の明細書はここに添付されているが、下記の額がチェックされている場合は、この限りでない：

the specification of which is attached hereto unless the following box is checked:

☐ _____ の日に出願され、
 この出願の米国出願番号または P C T 国際出願番号は、
 _____ であり、且つ
 _____ の日に補正された出願（該当する場合）

☒ was filed on September 19, 2000
as United States Application Number or
PCT International Application Number
PCT/JP00/06375 and was amended on
_____ (if applicable).

私は、上記の補正書によって補正された、特許請求範囲を含む上記明細書を検討し、且つ内容を理解していることをここに表明する。

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment referred to above.

私は、連邦規則法典第 37 編規則 1.56 に定義されている、特許性について重要な情報を開示する義務があることを認める。

I acknowledge the duty to disclose information which is material to patentability as defined in Title 37, Code of Federal Regulations, Section 1.56.

Burden Hour Statement: This form is estimated to take 0.4 hours to complete. Time will vary depending upon the need of the individual case. Any comments on the amount of time you are required to complete this form should be sent to Chief Information Officer, U.S. Patent and Trademark Office, Washington, DC 20231. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner of Patents and Trademarks, Washington, DC 20231.

Japanese Language Declaration (日本語宣言書)

私は、ここに、以下に記載した外国での特許出願または発明者証の出願、或いは米国以外の少なくとも一国を指定している米国法典第35編第365条(a)によるPCT国際出願について、同第119条(e)(d)項又は第365条(b)項に基づいて優先権を主張するとともに、優先権を主張する本出願の出願日より前の出願日を有する外国での特許出願または発明者証の出願、或いはPCT国際出願については、いかなる出願も、下記の枠内をチェックすることにより示した。

I hereby claim foreign priority under Title 35, United States Code, Section 119(a)-(d) or 365(b) of any foreign application(s) for patent or inventor's certificate, or 365(a) of any PCT International application which designated at least one country other than the United States listed below and have also identified below, by checking the box, any foreign application for patent or inventor's certificate, or PCT International application having a filing date before that of the application for which priority is claimed.

Prior Foreign Application(s)

外国での先行出願

266298/1999

(Number)
(番号)

Japan

(Country)
(国名)

20/09/1999

(Day/Month/Year Filed)
(出願日/月/年)

Priority

Claimed

☒ Yes ☐ No

357889/1999

(Number)
(番号)

Japan

(Country)
(国名)

16/12/1999

(Day/Month/Year Filed)
(出願日/月/年)☒ Yes ☐ No

126272/2000

(Number)
(番号)

Japan

(Country)
(国名)

20/04/2000

(Day/Month/Year Filed)
(出願日/月/年)☒ Yes ☐ No

私は、ここに、下記のいかなる米国仮特許出願についても、その米国法典第35編第119条(e)項の利益を主張する。

I hereby claim the benefit under Title 35, United States Code, Section 119(e) of any United States provisional application(s) listed below.

(Application No.)
(出願番号)(Filing Date)
(出願日)(Application No.)
(出願番号)(Filing Date)
(出願日)

私は、ここに、下記のいかなる米国出願についても、その米国法典第35編第120条に基づく利益を主張し、又米国を指定するいかなるPCT国際出願についても、その同第365条(c)に基づく利益を主張する。また、本出願の各特許請求の範囲の主題が、米国法典第35編第112条第1段に規定された態様で、先行する米国出願又はPCT国際出願に開示されていない場合においては、その先行出願の出願日と本国内出願日またはPCT国際出願日との間の期間中に入手された情報で、連邦規則法典第37編規則1.56に定義された特許性に関わる重要な情報について開示義務があることを承認する。

I hereby claim the benefit under Title 35, United States Code, Section 120 of any United States application(s), or 365(c) of any PCT International application designating the United States, listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States or PCT International application in the manner provided by the first paragraph of Title 35, United States Code Section 112, I acknowledge the duty to disclose information which is material to patentability as defined in Title 37, Code of Federal Regulations, Section 1.56 which became available between the filing date of the prior application and the national or PCT International filing date of application.

PCT/JP00/06375

September 19, 2000

(Application No.)
(出願番号)(Filing Date)
(出願日)

Pending

(Status: Patented, Pending, Abandoned)
(現況: 特許許可、係属中、放棄)(Application No.)
(出願番号)(Filing Date)
(出願日)(Status: Patented, Pending, Abandoned)
(現況: 特許許可、係属中、放棄)

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PTO/SB/106 (5-00)

Approved for use through 10/31/02. OMB 0651-0032

Patent and Trademark Office, U.S. DEPARTMENT OF COMMERCE

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委任状： 私は本出願を審査する手続を行い、且つ米国特許商標庁との全ての業務を遂行するために、記名された発明者として、下記の弁護士及び／または弁理士を任命する。(氏名及び登録番号を記載すること)

POWER OF ATTORNEY: As a named inventor, I hereby appoint the following attorney(s) and/or agent(s) to prosecute this application and transact all business in the Patent and Trademark Office connected therewith (list name and registration number).

Mark Chao, Reg. No. 37293; Elaine M. Ramesh, Reg. No. 43032

書類送付先

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Mark Chao, PhD. JD.

Intellectual Property Department

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唯一または第一発明者氏名

1-00

Full name of sole or first inventor

Kaneyoshi KATO

発明者の署名

日付

Inventor's signature

Date

Kaneyoshi Kato March 4, 2002

住所

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国籍

Citizenship

Japan

郵便の宛先

Post Office Address

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第二共同発明者がある場合、その氏名

2-00

Full name of second joint inventor, if any

Jun TERAUCHI

第二共同発明者の署名

日付

Second inventor's signature

Date

Jun Terauchi March 4, 2002

住所

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国籍

Citizenship

Japan

郵便の宛先

Post Office Address

same as above

(第三以下の共同発明者についても同様に記載し、署名をすること)

(Supply similar information and signature for third and subsequent joint inventors.)

Japanese Language Declaration

(日本語宣言書)

第三共同発明者がいる場合、その氏名	3-00	Full name of third joint inventor, if any	
		Masaaki MORI	
第三共同発明者の署名	日付	Third inventor's signature	Date
		<i>Masaaki Mori</i>	March 6, 2002
住所		Residence 8-5, Kasuga 3-chome,	
		Tsukuba-shi, Ibaraki 305-0821 Japan	
国籍		Citizenship	
		Japan	
郵便の宛先		Post Office Address	
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第四共同発明者がいる場合、その氏名	4-00	Full name of fourth joint inventor, if any	
		Nobuhiro SUZUKI	
第四共同発明者の署名	日付	Fourth inventor's signature	Date
		<i>Nobuhiro Suzuki</i>	March 4, 2002
住所		Residence 6-51, Fushiharacho,	
		Nishinomiya-shi, Hyogo 663-8031 Japan	
国籍		Citizenship	
		Japan	
郵便の宛先		Post Office Address	
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第五共同発明者がいる場合、その氏名	5-00	Full name of fifth joint inventor, if any	
		Yukio SHIMOMURA	
第五共同発明者の署名	日付	Fifth inventor's signature	Date
		<i>Yukio Shimomura</i>	March 6, 2002
住所		Residence 12-1-410, Matsushiro 3-chome,	
		Tsukuba-shi, Ibaraki 305-0035 Japan	
国籍		Citizenship	
		Japan	
郵便の宛先		Post Office Address	
		same as above	

Japanese Language Declaration
(日本語宣言書)

第六共同発明者がいる場合、その氏名	600	Full name of sixth joint inventor, if any	
		<u>Shiro TAKEKAWA</u>	
第六共同発明者の署名	日付	Sixth inventor's signature	Date
		<u>Shiro Takekawa</u>	March 4, 2002
住所		Residence	
		12-8-508, Miyanishicho, Nishinomiya-shi, Hyogo 662-0976 Japan	
国籍		Citizenship	
		Japan	
郵便の宛先		Post Office Address	
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第七共同発明者がいる場合、その氏名	700	Full name of seventh joint inventor, if any	
		<u>Yuji ISHIHARA</u>	
第七共同発明者の署名	日付	Seventh inventor's signature	Date
		<u>Yuji Ishihara</u>	March 4, 2002
住所		Residence	
		3-8, Yamada 3-chome, Itami-shi, Hyogo 664-0874 Japan	
国籍		Citizenship	
		Japan	
郵便の宛先		Post Office Address	
		same as above	

JPX

JPX

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3/11

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7/11

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Pro Gly Gly Ala Val Gly Cys Gly Ile Arg Leu Pro Asn Pro Asp Thr		
260	265	270
Asp Leu Tyr Trp Phe Thr Leu Tyr Gln Phe Phe Leu Ala Phe Ala Leu		
275	280	285
Pro Phe Val Val Ile Thr Ala Ala Tyr Val Arg Ile Leu Gln Arg MeT		
290	295	300
Thr Ser Ser Val Ala Pro Ala Ser Gln Arg Ser Ile Arg Leu Arg Thr		
305	310	315
Lys Arg Val Thr Arg Thr Ala Ile Ala Ile Cys Leu Val Phe Phe Val		
325	330	335
Cys Trp Ala Pro Tyr Tyr Val Leu Gln Leu Thr Gln Leu Ser Ile Ser		
340	345	350

Arg Pro Thr Leu Thr Phe Val Tyr Leu Tyr Asn Ala Ala Ile Ser Leu

355

360

365

Gly Tyr Ala Asn Ser Cys Leu Asn Pro Phe Val Tyr Ile Val Leu Cys

370

375

380

Glu Thr Phe Arg Lys Arg Leu Val Leu Ser Val Lys Pro Ala Ala Gln

385

390

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410

415

Thr Glu Ser Lys Gly Thr

420

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TCCACGGTCA TCTTCGCGGT CGTGAAGAAG TCCAAGCTGC ACTGGTGCAA CAACGTCCCC 240
GACATCTTCA TCATCAACCT CTCGGTAGTA GATCTCCTCT TTCTCCTGGG CATGCCCTTC 300
ATGATCCACC AGTCATGGG CAATGGGGTG TGGCACTTTG GGGAGACCAT GTGCACCCTC 360
ATCACGGCCA TGGATGCCAA TAGTCAGTTC ACCAGCACCT ACATCCTGAC CGCCATGGCC 420
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TGGCTGTATG CCAGACTCAT CCCCTTCCCA GGAGGTGCAG TGGGCTGCGG CATAAGCCTG 600
CCCAACCCAG AACTGACCT CTAAGGTTC ACCCTGTACC AGTTTTTCCT GGCCTTTGCC 660

CTGCCTTTTG TGGTCATCAC AGCCGCATAC GTGAGGATCC TGCAGCGCAT GACGTCCTCA 720
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CAGTTGTCCA TCAGCCGCCC GACCCTCACC TTTGTCTACT TATACAATGC GGCCATCAGC 900
TTGGGCTATG CCAACAGCTG CCTCAACCCC TTTGTGTACA TCGTGCTCTG TGAGACGTTT 960
CGCAAACGCT TGGTCCTGTC GGTGAAGCCT GCAGCCCAGG GGCAGCTTCG CGCTGTCAGC 1020
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<212> DNA

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CAAGCTGCAC TGGTGCAACA ACGTCCCCGA CATCTTCATC ATCAACCTCT CGGTAGTAGA 480
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GCACTTTGGG GAGACCATGT GCACCCTCAT CACGGCCATG GATGCCAATA GTCAGTTCAC 600
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CTCTTCCACG AAGTTCCGGA AGCCCTCTGT GGCCACCCTG GTGATCTGCC TCCTGTGGGC 720
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CCTGTACCAG TTTTTCCTGG CCTTTGCCCT GCCTTTTGTG GTCATCACAG CCGCATACGT 900
GAGGATCCTG CAGCGCATGA CGTCCTCAGT GGCCCCCGCC TCCCAGCGCA GCATCCGGCT 960
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